

UNIVERSITY of CALIFORNIA  
Santa Barbara

**High speed, manufacturable InP DHBTs and ICs with implanted collectors**

A dissertation submitted in partial satisfaction of the  
requirements for the degree of

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in

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by

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June 2006

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## PUBLICATIONS

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## Abstract

High speed, manufacturable InP DHBTs and ICs with implanted collectors

by

Navin Parthasarathy

Novel HBT technologies have been designed and developed to enable continued scaling of InP based double heterojunction bipolar transistors(DHBTs), for their use in high speed, low power digital logic and mixed-signal systems. Significant reduction in parasitics have been achieved by the use of implanted subcollector and pedestal for independent collector scaling .

The base collector capacitance  $C_{cb}$ , contributes significantly to analog bandwidth and digital delay and must be reduced as DHBTs are scaled. Two different types of InP double heterojunction bipolar transistors have been designed, fabricated and characterized employing Fe and selective Si implants to decrease  $C_{cb}$ . The first type, the selectively implanted subcollector DHBT, has a shallow Fe implant to compensate regrowth interface charge and a Si implant to form the subcollector. This selective implant eliminates the parasitic base collector capacitance associated with the base access pad area. These devices exhibit 361 GHz  $f_{\tau}$  and 404 GHz  $f_{max}$ . A second device, the implanted pedestal-subcollector DHBTs allows for further reduction of  $C_{cb}$ . It has a thicker Fe implanted semi-insulating layer for reduced  $C_{cb}$  in the ex-

trinsic collector-base junction, a patterned buried subcollector formed by a deep Si implant and an collector pedestal created by a second Si implant. These InP pedestal HBTs have 352 GHz  $f_\tau$  and 403 GHz  $f_{max}$ . The implanted pedestal-subcollector DHBTs have a DC current gain  $\beta \approx 35$  and  $BV_{CBO} \approx 7.8$  V. In addition to the compensation by the Fe implant of charge at the epitaxial growth interface, these two processes provide the following enhancements: elimination of  $C_{cb}$  in the base interconnect pad area, a single MBE growth and increased wafer planarity. In this first demonstration of these two processes, the  $f_\tau$  and  $f_{max}$  are the highest reported for DHBTs with implanted collectors. Using these implanted collector processes, record low power delay products have been obtained for CML dividers operating at over 61 GHz with an operating power of less than 27mW. High speed CML static frequency dividers clocking to 135 GHz have been measured. ECL dividers utilizing implanted collectors have a self oscillation frequency of 96 GHz. There is a 20 % improvement in logic speed using this technology.

For high speed logic circuits, emitter resistance is an important parameter to be reduced. Highly scaled emitter junctions with simultaneously large emitter contacts, and thick extrinsic bases have been demonstrated with an emitter regrowth technology. These have attained 280 GHz  $f_\tau$ . In conjunction with the implanted collector process, these InP DHBTs can attain a bandwidth of over 1 THz.



To  
my Mother, Father and Ammu

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# 1

## Introduction

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HIGH performance, manufacturable transistors are required for significant improvements in the bandwidth, dynamic range, power consumption and integration level of mixed-signal ICs used in military radar and communications transmitters. Large integration scales,  $\approx 20,000$  transistors, will be needed to realize complex signal processing functions, and power per gate must be accordingly reduced to achieve acceptable thermal densities for reliable device operation. Future microwave DACs and DDFS ICs will operate at 100-200 GHz clock speeds and will contain  $10^4 - 10^5$  transistors. This requires a breakthrough in combined IC speed and integration scales.

Despite tremendous advancements in CMOS technology, bipolar transistors remain competitive owing to larger breakdown voltages obtainable and more tolerant lithographic dimensions at a given bandwidth. SiGe and InP heterojunction transistors (HBTs) are the main competing high speed IC technologies. InP heterojunction transistors benefit from high carrier mobilities and saturated velocities and are compatible for integration with 1.3 - 1.5  $\mu\text{m}$  optoelectronic components such as lasers and photodetectors [1]. SiGe HBTs have smaller junction dimensions and smaller extrinsic parasitics due to the maturity of the advanced silicon processes. Consequently, digital circuit speed in SiGe and InP has been comparable [2], with SiGe offering higher integration scales. Further scaling of InP HBTs is therefore important, and with the adoption of advanced SiGe-like fabrication processes, large potential improvements in speed and yield of InP HBTs are possible. InP HBT processes utilizing  $\approx 500$  nm width emitter junctions ( $W_e$ ) and 500 nm width base contacts ( $W_b$ ) have demonstrated 350 GHz  $f_\tau$ , 400 GHz  $f_{max}$  and 150 GHz static frequency dividers [3, 4].

For  $\approx 2:1$  improvement in bandwidth, the collector layer must be thinned 2:1 and the base thinned  $\sqrt{2}:1$ . The operating current density must be increased 4:1 and the emitter resistance per unit emitter junction area reduced 4:1. Emitter and collector junction widths must both decrease 4:1. If the base contacts lie above

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the active collector-base junction, the contact width and the contact resistivity must both decrease 4:1 [5]. In addition to difficulties in reducing contact resistivity, very narrow base contacts present challenges in process design for high yield fabrication and present significant bulk metal resistance along the length of the contact. Further, the parasitic base interconnect pad capacitance becomes a significant fraction of the total in small-area HBTs used in low-power logic.

This work describes InP DHBTs that address the above scaling issues and suggests a restructuring of the III-V fabrication scaling process to closely follow the Si/SiGe techniques. Fabrication steps have been developed aimed at improving the performance and manufacturability of a submicron InP DHBT device. The device epitaxy has been tailored to support a submicron process, with device parameters that are optimized for high digital logic speeds and not necessarily traditional transistor figures-of-merit ( $f_\tau$ ,  $f_{max}$ ). Work has been presented on two unique device topologies.

The first topology is a selectively implanted subcollector and pedestal HBT technology. Scaling of the collector base junction is achieved by the use of Fe and selective Si implants. DHBTs with implanted subcollector DHBTs are fabricated where the parasitic capacitance associated with the area of the base access pad  $C_{cb,pad}$ , is eliminated. A reliable method for compensating the charge associated with the InP growth interface is also demonstrated. Further reduction in extrinsic collector base capacitance  $C_{cb}$ , is achieved by the addition of a selectively implanted pedestal in the second device topology. InP DHBTs employing a buried  $N^{++}$  sub-collector and  $N^+$  collector pedestal formed fully by ion implantation are demonstrated. This novel DHBT structure provides the following enhancements over the existing pedestal InP DHBTs [6, 7]: reduced extrinsic  $C_{cb}$  associated with the pedestal, compensation of charge at the epitaxial growth interface, elimination of  $C_{cb}$  in the base interconnect pad area, a single MBE growth and increased wafer planarity and hence potentially improved yield in the fabrication of large circuits. In this first demonstration of the fully implanted pedestal-subcollector DHBTs, the  $f_\tau$  and  $f_{max}$  are a record for transistors with implanted collectors.

This thesis is logically structured to explain the steps taken towards a SiGe-like highly scaled, manufacturable InP HBT. Chapter 2 discusses the basic theory and design of an InP based DHBT. Scaling laws for increasing HBT and digital IC bandwidth, and physical limits to scaling are presented. Chapter 3 presents the theory of ion implantation in InP and discusses the design of Silicon (Si) and Iron (Fe) implants in InP. The results pertaining to Si and Fe implants and high temperature annealing of InP are presented. Chapter 4 may be considered as two parts. In the first half, device results are presented for a DHBT with implanted subcollectors formed by a single Si implant. The effect of growth interface charge is shown and an approach to eliminate this is proposed and developed. The second half, disc-

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cuses the fabrication and device results of implanted subcollector DHBTs with two ion implants are discussed. Chapter 5 describes in depth the design and performance of the implanted pedestal-subcollector DHBT. Chapter 6 summarises the research effort and outlines directions for future work.

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# 2

## InP DHBT Theory and Design

---

ADVANCES in the design and fabrication of InP heterojunction bipolar transistors have been made at UCSB since 1994. In this chapter, the InP HBT layer structure and HBT scaling laws are presented. The transit times, resistances and capacitances are examined. The various transistor figures-of-merit are described. The delays associated with a digital latch, regularly employed as retiming elements and decision circuits is derived. Lastly, device modeling and scaling limits are discussed.

### 2.1 Emitter

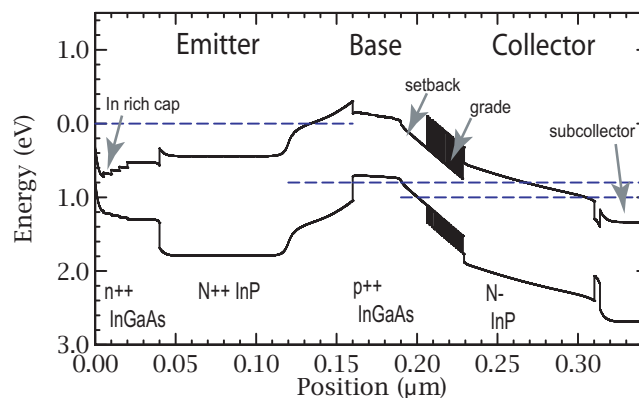


Figure 2.1: Energy band diagram of a typical HBT with  $V_{be} = 0.9\text{V}$  and  $V_{cb} = 0.2\text{V}$

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The band diagram of a typical NpN InP based DHBT is shown in Fig. 2.1. Under normal operation, the emitter base junction is forward biased. This lowers the emitter base potential and injects electrons into the base. As seen in Fig. 2.2, the emitter layer stack consists of very highly doped  $In_xGa_{1-x}As$  used as the contact

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Thickness (nm)	Semiconductor Composition	Doping ( $\text{cm}^{-3}$ )	Description
5	$\text{In}_{0.85}\text{Ga}_{0.15}\text{As}$	$> 5 \cdot 10^{19}:\text{Si}$	Emitter cap
15	$\text{In}_x\text{Ga}_{1-x}\text{As}$	$4 \cdot 10^{19}:\text{Si}$	Emitter cap
20	$\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$	$3 \cdot 10^{19}:\text{Si}$	Emitter cap
80	InP	$3 \cdot 10^{19}:\text{Si}$	Emitter
10	InP	$8 \cdot 10^{17}:\text{Si}$	Emitter
40	InP	$5 \cdot 10^{17}:\text{Si}$	Emitter
30	InGaAs	$4 - 7 \cdot 10^{19}:\text{C}$	Base
15	$\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$	$3.25 \cdot 10^{16}:\text{Si}$	Setback
24	InGaAs / InAlAs	$3.25 \cdot 10^{16}:\text{Si}$	B-C grade
3	InP	$2.75 \cdot 10^{18}:\text{Si}$	Delta doping
78	InP	$3.25 \cdot 10^{16}:\text{Si}$	Collector
5	InP	$1.5 \cdot 10^{19}:\text{Si}$	Sub-collector
6.5	$\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$	$2 \cdot 10^{19}:\text{Si}$	Sub-collector
300	InP	$2 \cdot 10^{19}:\text{Si}$	Sub-collector
Substrate	Semi-Insulating InP		

Figure 2.2: Layer structure of a typical UCSB mesa-DHBT

layer and a  $N^-$  InP emitter. The InP-InGaAs base emitter heterojunction can be abrupt or graded. The DC current gain of a graded junction is higher than that of an abrupt junction since the hole barrier is larger. The ratio of emitter current between an abrupt and a graded emitter junction is

$$\frac{I_{E,abrupt}}{I_{E,graded}} \simeq \exp\left(-\frac{\Delta E_c}{kT}\right) \quad (2.1.1)$$

where  $\Delta E_c$  is the conduction band discontinuity between InP and  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  of the base. However grading presents many challenges and the base-emitter grade design is not straightforward for the forward biased junction [1]. It can be argued that for an abrupt emitter-base junction, the electrons that surmount the barrier are injected with a substantial kinetic energy roughly equal to  $\Delta E_c$ . This kinetic energy corresponds to a high velocity which influences the base and collector transit times [2]. Besides the choice between a graded and abrupt base-emitter heterojunction, the 3 parameters about the emitter design are the InP emitter layer thickness ( $X_E$ ), doping level ( $N_E$ ) and the choice of emitter cap.  $X_E$  and  $N_E$  influence the base-



## CHAPTER 2. INP DHBT THEORY AND DESIGN

emitter junction capacitance.  $C_{je}$  per unit area is given by

$$C_{je} = \frac{\epsilon_{InP}}{X_{dep,E}} \simeq \frac{\epsilon_{InP}}{\sqrt{\frac{2\epsilon_{InP}}{qN_E}(\phi_{BE} - V_{BE})}} \quad (2.1.2)$$

$C_{je}$  is not purely a depletion capacitance and making  $X_{dep,E}$  (or  $X_E$ ) large fails in the limit of large depletion thicknesses. The emitter base junction capacitance  $C_{je}$  can also be given by,

$$C_{je} = C_{je1} + C_{je2} = \frac{\epsilon_{InP}L_eW_e}{X_{dep,E}} + \kappa X_{dep,E}T_bI_c \quad (2.1.3)$$

where  $T_b$  is the base thickness and  $\kappa$  is a constant. If the current density is increased, while maintaining constant  $X_{dep,E}$ , the stored mobile electron charge is increased. It can be shown that the ideality factor  $\eta$  is,

$$\eta = 1 + \frac{1}{q} \frac{\partial(\Delta E_{fn,eb})}{\partial V_{be}} \quad (2.1.4)$$

where  $\Delta E_{fn,eb}$  is the change in quasi-Fermi energy due to recombination at the base-edge of the emitter. To maintain a high current density without significant voltage drop in the base-emitter depletion region, a high electron density is to be maintained. However, to obtain a low ideality factor, the electron concentration in the depletion region must be kept high while keeping the thickness of the depletion region small. The emitter charging time  $\tau_e = (kT/\eta q I_c) \cdot C_{je}$  is one of the major delay components determining  $f_\tau$  and maximum digital logic speed [3].

The delay term  $R_{ex} C_{cb}$  is a major limit to HBT scaling and  $R_{ex}$  degrades the digital noise margin. The emitter layer structure consists of a heavily doped and narrow bandgap contact layer (cap), a highly doped  $N^{++}$  wide bandgap emitter and a low doped  $N^-$  emitter of thickness  $X_E$  and doping  $N_E$ . If the heterointerfaces are designed properly to avoid conduction band barriers between layers, and if the thicknesses of the  $N^{++}$  layers are not significantly larger than  $X_E$ , the parasitic emitter resistance is determined mostly by the contact resistance between the metal and the semiconductor and the resistance in the undepleted portion of the  $N^-$  emitter. The emitter resistance for a standard mesa DHBT is then given by

$$R_{ex} \simeq \frac{\rho_{c,e}}{L_e W_{ec}} \quad (2.1.5)$$

where  $\rho_{c,e}$  is the emitter specific contact resistivity and  $\rho_{e1}$  is the bulk resistivity of the  $N^-$  emitter layer and the other dimensions are as shown in Fig. ?? . For submicron junctions, the junction width  $W_{ej}$  is smaller than the contact width due

## CHAPTER 2. INP DHBT THEORY AND DESIGN

to the lateral undercutting during the wet etching of the emitter. Surface depletion of the  $N^-$  layer further reduces the width of the electrically active junction. An undoped emitter is reported [4] to provide a higher  $f_\tau$  at a given current density  $J_c$ . However, low doping leads to higher emitter charging times at high current densities.

### 2.1.1 Emitter Resistance

The emitter resistance is primarily determined by the ohmic contact resistance  $\rho_c$ . The ohmic resistivity  $\rho_c$  is,

$$\rho_c \propto \exp \left[ \frac{\Phi_B}{N^{\frac{1}{2}}} \right] \quad (2.1.5)$$

The resistivity is proportional to the barrier height  $\Phi_B$  and the doping of the emitter cap [5]. The resistivity can be reduced by using a highly doped, low bandgap semiconductor as the emitter cap. For the DHBTs described here, the emitter cap is  $\text{In}_{0.85}\text{Ga}_{0.15}\text{As}$  which has a lower band gap than  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$ . These are doped at over  $3 \times 10^{19} \text{ cm}^{-3}$ . The maximum doping is limited by the solid solubility of Si dopants in  $\text{In}_{0.85}\text{Ga}_{0.15}\text{As}$ . The thickness of the emitter cap is 40 nm. This is limited by the Matthews-Blakeslee critical limit for strained  $\text{In}_{0.85}\text{Ga}_{0.15}\text{As}$  growth. Furthermore, the cap is kept thin to prevent excessive undercut of the  $\text{In}_{1-x}\text{Ga}_x\text{As}$  layers during the emitter wet etch.

## 2.2 Base

The base-emitter turn on voltage is approximately the built in potential and for the abrupt emitter-base junction used in the designs in this thesis is given by,

$$\phi_{bi} = \frac{E_{gb} + \Delta E_c - \phi_p - \phi_n}{q} \quad (2.2.0)$$

$E_{gb}$  is the bandgap of of the base layer. Ignoring high doping effects,  $E_g$  is  $\approx 0.76$  eV for  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$ ,  $\Delta E_c$  is the conduction band offset to InP  $\approx 0.26$  eV.  $\phi_p$  is the hole quasi Fermi level in the base with respect to the valence band and  $\phi_n$  is the electron quasi Fermi level in the emitter with respect to the conduction band. For non-degenerately doped semiconductors, these electron and hole quasi Fermi levels are described by,

$$\phi_n = E_c - E_{fn} \quad (2.2.1)$$

$$\phi_p = E_{fp} - E_v \quad (2.2.2)$$

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The  $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  base is usually between 20-100nm and doped well over  $10^{19}\text{cm}^{-3}$ . The electron and hole currents in the base are,

$$J_n = \mu_n \frac{d}{dx} \phi_n \quad (2.2.3)$$

$$J_p = \mu_p \frac{d}{dx} \phi_p \quad (2.2.4)$$

The hole current should be very small  $\approx 0$  in a well designed bipolar transistor. The following generalized Moll-Ross current relation for a bipolar transistor with a non-uniform base is derived by Kroemer [6] as

$$J_n \approx - \frac{q \cdot \exp\left(\frac{qV_{be}}{kT}\right)}{\int_{base} [p/D_n \cdot n_{ib}^2] dx} \quad (2.2.4)$$

Here,  $D_n$  is the electron diffusivity in the base and  $n_{ib}$  is the intrinsic carrier concentration in the base.

The time it takes for an electron entering from the emitter to traverse across the base is the base transit time and can be expressed as,

$$\tau_b = \frac{Q_B}{I_n} = \frac{\int_0^{T_b} q \cdot n(x) dx}{J_n} \quad (2.2.4)$$

where  $T_b$  is the base thickness. When the electron transport is purely diffusive, the transit time is classically given by,

$$\tau_b = \frac{T_b^2}{2D_n} \quad (2.2.4)$$

This transit time calculation assumes uniform composition and doping in the base. To reduce  $\tau_b$ , an electric field can be established in the base so that electron transport is aided by the drift field [7]. If the grading of the base conduction band is linear, Equ. 2.2 is rewritten as,

$$\tau_b = \frac{T_b^2}{D_n} \left( \frac{kT}{\Delta E} \right) \left[ 1 - \frac{kT}{\Delta E} \left( 1 - \exp^{-\Delta E/kT} \right) \right] \quad (2.2.4)$$

where  $\Delta E$  is the potential due to grading across the base. The grading may be done compositionally to vary the base bandgap or by varying the base doping. The HBTs presented in this work employ a doping grade producing a  $\Delta E \sim 50$  meV. This in turn reduces  $\tau_b$  by  $\sim 50\%$  compared to an ungraded base. For transistors with very

## CHAPTER 2. INP DHBT THEORY AND DESIGN

thin bases operating at high current densities, Shockley's boundary conditions cases to be valid

$$\tau_b = \frac{T_b^2}{D_n} \left( \frac{kT}{\Delta E} \right) \left[ 1 - \frac{kT}{\Delta E} \left( 1 - \exp^{-\Delta E/kT} \right) \right] + \frac{T_b}{v_{exit}} \left( \frac{kT}{\Delta E} \right) \left( 1 - \exp^{-\Delta E/kT} \right) \quad (2.2.4)$$

The  $T_b/v_{exit}$  contribution is usually ignored by assuming that the electron concentration at the collector side of the base is zero. For a thin base this assumption is not valid and the correction term accounts for the finite electron concentration that exits the base. For bases thicker than 20nm, the base transit time  $\tau_b \sim T_b^2$ , while for bases thinner than 20nm, the base transit time  $\tau_b \sim T_b$ .

The base current components in a well designed HBT consists of bulk recombination current, surface recombination current, emitter-base space charge recombination current. These components need to be minimised to increase the current gain of the transistor. The surface recombination current is directly proportional to the emitter periphery. Unlike the GaAs surface, the free  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  has a much lower surface recombination velocity of  $\sim 1 \times 10^3 \text{ cm/s}$ . Further, BCB is used for device passivation and reduces the surface leakage current, compared to polyimide or  $\text{Si}_x\text{N}_y$  [8]. The base current is dominated by Auger recombination in the bulk. The Auger recombination rate is given as

$$U_A = C(n+p)(np - n_i^2) \approx (CN_A^2) \cdot \Delta n = \frac{\Delta n}{\tau_r} \quad (2.2.4)$$

where C is the Auger recombination co-efficient and  $\Delta n$  is the excess electron concentration in the base and  $\tau_r$  is called the recombination lifetime and is a material parameter. The current gain can now be expressed as

$$\beta = \frac{\tau_n}{\tau_r} \quad (2.2.4)$$

The current gain is thus inversely proportional to the base thickness and the base doping. Reducing the base thickness and base doping however increases the base resistance and impacts the  $f_{max}$  of the bipolar transistor. The base resistance for a two sided base contact is given by,

$$R_{bb} = R_{bb,contact} + R_{bb,gap} + R_{bb,spread} \quad (2.2.5)$$

$$R_{bb,contact} = \frac{\sqrt{\rho_c \cdot R_{sh,b}}}{2L_E} \coth \frac{W_B}{L_T} \text{ where, } L_T = \sqrt{\rho_c/R_{sh,b}}$$

$$R_{b,gap} = R_{sh,b} \frac{W_{gap}}{2L_E} \text{ and } R_{b,spread} = R_{sh,b} \frac{W_E}{12L_E}$$

$$R_{bb} = \frac{1}{2} \frac{R_{sh,b}}{L_e} \left[ L_t \coth \frac{W_b}{L_t} + W_{gap} + \frac{W_e}{6} \right] \quad (2.2.4)$$

## CHAPTER 2. INP DHBT THEORY AND DESIGN

where  $\rho_c$  ( $\Omega \cdot \mu\text{m}^2$ ) and  $R_{sh,b}$  ( $\Omega/\square$ ) are the specific contact resistivity and sheet resistance of the base, and  $L_T$  is called the ohmic transfer length. The base contact resistance is  $\propto L_T$ . Therefore, scaling down the base-collector junction laterally increases the base resistance exponentially as seen in Eqn. 2.2.4. The base resistance is usually higher than the emitter and collector resistances due to the heavier effective mass and lower mobility of the hole.

When current is flowing through the device, an important reduction of extracted value of  $R_{bb}$  is observed as seen on Fig. 2.10 in §???. This decrease of  $R_{bb}$  is due to decrease in  $R_{bb,spread}$ , attributed to electron injection in intrinsic base [?]. If average electron velocity in the is about  $3 \times 10^7$  cm/s, an emitter current density of  $2$  mA/ $\mu\text{m}^2$  induces an electron concentration of  $4 \times 10^{16}$   $\rho_c$ . This modifies the resistivity in the intrinsic base, due to the high electron mobility in  $p^{++}$  In<sub>0.53</sub>Ga<sub>0.47</sub>As of  $\sim 3000$   $\text{cm}^2/\text{V} \cdot \text{s}$ , compared to the hole mobility of  $\approx 40$   $\text{cm}^2/\text{V} \cdot \text{s}$ .

### 2.3 Collector

The collector design is critical to the performance of high speed transistors. There are breakdown advantages when the collector layer is made of a widegap material. In InP/InGaAs DHBTs, there is a discontinuity in the conduction band between the In<sub>0.53</sub>Ga<sub>0.47</sub>As base and the InP collector layers. In order to prevent blocking of current, this discontinuity is removed by grading the energy gap from the base to the collector. The collector layer is thus a composite structure consisting of a In<sub>0.53</sub>Ga<sub>0.47</sub>As setback, a In<sub>0.53</sub>Ga<sub>0.47</sub>As- InP grade and wide bandgap InP collector as seen in the energy band diagram in Fig. ??. The base-collector grading can be accomplished in one of two ways

- In<sub>1-x</sub>Ga<sub>x</sub>As<sub>y</sub>P<sub>1-y</sub>quarternary lattice matched to InP. By varying the ratio of the Group V element (As, P), from  $y=1 \rightarrow y=0$ , the energy band is progressively graded from In<sub>0.53</sub>Ga<sub>0.47</sub>As to InP [9]. This is particularly convenient for MOCVD growth.
- InGaAs/InAlAs super-lattice grade with intermediate effective bandgap [10]. A chirped superlattice of In<sub>0.53</sub>Ga<sub>0.47</sub>As and In<sub>0.52</sub>Al<sub>0.48</sub>As is form results in delocalised electron states resulting in an effective bandgap. The periodicity has to be kept small, compared to the electron wavelength, to prevent electron reflection. In all the designs described in this thesis, the super lattice period is 1.5nm.

The electric field due to the grade, given by  $\Delta E_c/T_{grade}$ , opposes the built-in field. Two  $\delta$  doping layers at the end of the grading length  $T_{grade}$ , acceptor at the base end

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and donor at the collector end forms a dipole that cancels out the electrostatic field due to the grade [11]. The sheet charge density of this doping layer is given by

$$N_\delta T_\delta = \frac{\varepsilon_o \varepsilon_r \Delta E_c}{q^2 \cdot T_{grade}} \quad (2.3.0)$$

where  $N_\delta$  and  $T_\delta$  are the doping concentration and thickness of the  $\delta$  layer. A setback layer of N-type  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  is usually inserted between the base and the grade. This layer serves to ensure that have electrons have sufficient kinetic energy before entering the grade [10]. The potential drop in the setback layer,  $\Delta\phi_{setback}$ , can be a significant fraction of  $\phi_{bi}$  which could result in lower breakdown. A rule of thumb is to design the setback such that  $\Delta\phi_{setback} = \Delta E_c$ . The rest of the collector consists of wide bandgap InP. The doping in the collector is chosen so that the collector is fully depleted at zero bias. To ensure full depletion in the collector at zero bias,

$$\phi_{bi} \geq \frac{qN_c(T_{setback} + T_{grade})^2}{2\varepsilon_o\varepsilon_r} + \frac{qN_\delta T_\delta(T_{setback} + T_{grade})}{\varepsilon_o\varepsilon_r} + \frac{qN_c T_{InP}^2}{2\varepsilon_o\varepsilon_r} - \frac{T_{grade}\Delta E_c}{q} \quad (2.3.0)$$

where  $T_{setback}$ ,  $T_{grade}$  and  $T_{InP}$  are the thicknesses of the setback, grade and InP layers respectively.  $N_c$  is the doping of the collector. From this expression the maximum allowable collector doping to ensure full collector depletion is,

$$N_{c,max} \approx \frac{1}{T_{InP}^2} \left[ \frac{2\varepsilon_o\varepsilon_r}{q} (\phi_{bi} - \Delta\phi_{setback} - \Delta\phi_{grade}) \right]$$

where  $\Delta\phi_{setback}$  and  $\Delta\phi_{grade}$  are the potential drops in the setback and grade layers. The potential dipole reduces the maximum collector doping significantly.

The transit time for an electron to through the collector is given by

$$\tau_c = \int_0^{T_c} \frac{1 - x/T_c}{v(x)} dx \equiv \frac{T_c}{2v_{eff}}$$

where  $\tau_c$  is the collector transit time,  $v(x)$  and  $v_{eff}$  are the position-dependent and effective electron velocities in the collector drift region.

The subcollector layer is 300 nm of Si doped  $N^{++}$  InP. The subcollector is designed to reduce the collector resistance. The expression for collector resistance of a standard mesa DHBT, for a two sided collector contact is,

$$R_c = R_{c,contact} + R_{c,gap} + R_{c,spread} \quad (2.3.0)$$

$$R_c = \frac{1}{2} \frac{R_{sh,s}}{L_e} \left[ L_t + W_{c,gap} + \frac{W_{cb}}{6} \right]$$

## CHAPTER 2. INP DHBT THEORY AND DESIGN

where  $\rho_c$  ( $\Omega \cdot \mu\text{m}^2$ ) and  $R_{sh,s}$  ( $\Omega/\square$ ) are the specific contact resistivity and sheet resistance of the sub-collector,  $W_{cb}$  is the width of the collector mesa,  $W_{c,gap}$  is the spacing between the collector mesa and contact,  $L_e$  is the emitter junction length, and  $L_t$  is the ohmic transfer length equal to  $\sqrt{\rho_c/\rho_s}$ .  $R_{gap}$  and  $R_{c,spread}$  are reduced by decreasing the sheet resistance of the subcollector which can be expressed as

$$R_{sh,s} = \frac{1}{q\mu_n \cdot N_{sc} \cdot T_{sc}} \quad (2.3.0)$$

where  $T_{sc}$  is the thickness of the subcollector layer,  $\mu_n$  is the mobility of the majority electrons, in  $N^{++}$  InP,  $\mu_n \propto N^{-0.2}$  [12]. Si is an amphoteric dopant in InP and above certain doping levels ( $\sim 4\text{E}19 \text{ cm}^{-3}$ ) starts to self compensate. The doping is chosen to obtain the lowest possible resistivity in the  $N^{++}$  layer. From the above expression, increasing the the subcollector thickness reduces the sheet resistance but this  $T_{sc}^{-1}$  decrease is insignificant beyond  $\approx 0.5 \mu\text{m}$ . There is a practical concern for thick subcollectors in a standard triple mesa DHBT where device isolation is done by etching through the subcollector to the substrate. A thick subcollector thus results in a highly non-planar device which can compromise yield. A major portion of this thesis addresses these issues through the use of implanted subcollectors described in Chapter ?? . To reduce the contact resistance, a thin layer of highly doped  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  is used as the contact layer in standard triple mesa DHBTs [13]. Ternary materials have poor thermal conductivity (0.05 W/cm-K for  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  vs. 0.88 W/cm-K for InP) due to phonon Rayleigh scattering from the alloy disorder. This layer is kept thin to minimise the thermal resistance [14]. As will be described in detail in Chapter ?? , in the implanted DHBTs this InGaAs layer is undoped and designed to be 3.5 nm and is used as an etch-stop layer. The collector contact in these DHBTs is made to  $N^{++}$  InP.

### 2.3.1 Base-collector depletion capacitance, $C_{cb}$

The depleted space charge in the collector can be modeled as a parallel plate capacitance [15]. The collector is designed to be fully depleted at zero bias. This ensures a minimal variance in collector base capacitance as increased reverse bias is applied. For the mesa HBT structure shown in Fig. 2.3, the four components of the collector-base capacitance are,

$$C_{cb} = C_{cb,ex} + C_{cb,gap} + C_{cb,i} + C_{cb,pad} \quad (2.3.0)$$

$$C_{cb,ex} = 2 \frac{\varepsilon_o \varepsilon_r L_e W_b}{T_c}, \quad C_{cb,gap} = 2 \frac{\varepsilon_o \varepsilon_r L_e W_{gap}}{T_c}, \quad C_{cb,i} = \frac{\varepsilon_o \varepsilon_r L_e W_e}{T_c}, \quad C_{cb,pad} = 2 \frac{\varepsilon_o \varepsilon_r A_{pad}}{T_c}$$

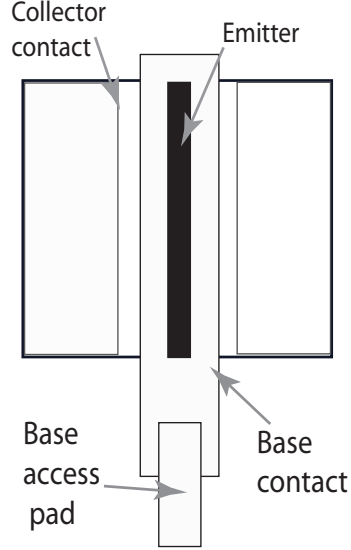


Figure 2.3: Top view of the mesa HBT showing the metal contacts

, where  $W_b$  is the width of the base metal-semiconductor junction,  $W_{b,gap}$  is the spacing between the base contact and emitter mesa, and  $W_e$  and  $L_e$  are the width and length of the emitter junction.  $A_{pad}$  is the base access pad as seen in Fig. 2.3. A major portion of this thesis concerns the reduction of the collector-base capacitance. The collector base capacitance can be modeled as

$$C_{cb} = -\frac{\partial Q_{base}}{\partial V_{cb}} \quad (2.3.0)$$

But the transit time  $\tau_f$ ,

$$\tau_f = \tau_c + \tau_b \equiv \frac{\partial Q_{base}}{\partial I_c} \quad (2.3.0)$$

Therefore from Eqns. 2.3.1 and 2.3.1,

$$\frac{\partial C_{cb}}{\partial I_c} = -\frac{\partial \tau_f}{\partial V_{cb}} \quad (2.3.0)$$

If  $\frac{\partial \tau_f}{\partial V_{cb}} > 0$ , then  $\frac{\partial C_{cb}}{\partial I_c} < 0$ . This is called collector velocity modulation.

## 2.4 Figures of merit

Various figures of merit are used to describe the transistor.



### 2.4.1 Current-gain cutoff frequency $f_\tau$

The current gain cut-off frequency for a bipolar transistor is given by [16],

$$\frac{1}{2\pi f_\tau} = \tau_c + \tau_b + C_{cb} \cdot (R_{ex} + R_c) + \frac{\eta kT}{qI_e}(C_{cb} + C_{je}) \quad (2.4.0)$$

where  $\tau_c$  and  $\tau_b$  are the collector and base transit times,  $C_{cb}$  and  $C_{je}$  are the depletion capacitances for the collector and emitter,  $R_c$  and  $R_{ex}$  are the series resistances of the collector and emitter, and  $(\eta kT/qI_e)^{-1}$  is the transconductance of the HBT. For a typical mesa DHBT with 120nm collector, 30nm base, emitter junction =  $0.65 \times 4.3 \mu m^2$ ,  $\rho_{c,C} = \rho_{c,E} = 10 \Omega \cdot \mu m^2$ , base mesa width =  $1.3 \mu m$ ,  $\tau_c = 0.188 ps$ ,  $\tau_b = 0.104 ps$ ,  $C_{cb} \cdot (R_{ex} + R_c) = 0.044 ps$ ,  $\frac{\eta kT}{qI_e}(C_{cb} + C_{je}) = 0.030 ps$ . The collector and base transit delay are seen to be dominant for the HBT described. As the devices are scaled vertically, these delays decrease. However the time constants associated with charging the base-collector capacitance starts to dominate. Thus reduction of base-collector capacitance gains significance as devices are scaled.

### 2.4.2 Power-gain cutoff frequency $f_{max}$

The HBT maximum oscillation (unity power-gain) frequency is,

$$f_{max} = \sqrt{\frac{f_\tau}{8\pi(R_{bb}C_{cb})_{eff}}} \quad (2.4.0)$$

dependent upon the HBT  $f_\tau$  and a time constant  $(R_{bb}C_{cb})_{eff}$  that includes the effects of the distributed base-collector network [17]. Each component of  $C_{cb}$  is charged through a certain portion of the base resistance  $R_{bb}$ . Utilizing the definitions for  $C_{cb}$  and  $R_{bb}$  from Equ. 2.3.1 and 2.2.4 and assuming little undercut of the base-collector mesa,

$$(R_{bb}C_{cb})_{eff} \approx C_{cb,i}R_{bb} + C_{cb,gap}(R_{b,cont} + R_{b,gap}/2) + C_{cb,ex}R_{b,cont,0} + C_{pad}R_{b,cont,pad} \quad (2.4.0)$$

where the collector-base capacitance underneath the emitter stripe  $C_{cb,i}$  is charged through the entire base resistance, and the gap capacitance  $C_{cb,gap}$  between the emitter mesa and base contact is charged through  $(R_{b,cont} + R_{b,gap}/2)$ . The extrinsic collector-base capacitance  $C_{cb,ex}$  underneath the base contact is charged by currents traversing vertically through the contact above it, having a resistance  $R_{b,cont,0} = \rho_c/L_e W_b$ . The pad capacitance is charged by the vertical contact resistance associated with the base access pad,  $R_{b,cont,pad} = \rho_c/A_{pad}$ ). For the sake of simplicity, Eqn. 2.4.2 for  $f_{max}$  ignores the effect of the collector and emitter series resistances since  $R_{bb}$  is usually much larger than  $R_c$  and  $R_{ex}$ . Power is dissipated in these resistances and their effect must be included especially if these resistances are  $\sim R_{bb}$ .

### 2.4.3 Maximum current density $J_{kirk}$

As the collector current density is varied, the injected electrons screen the background doping and modify the electric field profile in the collector. To account for this injected charge, Poisson equation can be written,

$$-\frac{d^2\phi}{dx^2} = \frac{dE}{dx} = \frac{1}{\varepsilon_o\varepsilon_r} \left[ qN_c - \frac{J(x)}{v(x)} \right] \quad (2.4.0)$$

where  $\varepsilon_o\varepsilon_r$  is the dielectric constant,  $N_c$  the collector doping,  $J(x)$  the collector current density, and  $v(x)$  the electron velocity in the collector. Integrating Equ. 2.4.3 to solve for the electric field  $E(x)$ ,

$$E(x) = \frac{1}{\varepsilon_o\varepsilon_r} \int_0^x qN_c - \frac{J(x)}{v(x)} dx \quad (2.4.0)$$

Assuming  $J(x)$  and  $v(x)$  to be constant and integrating the electric field over the collector to obtain the potential,

$$\phi_{bi} + V_{cb} \geq \frac{T_c^2}{2\varepsilon_o\varepsilon_r} \left[ qN_c - \frac{J_e}{v_{eff}} \right] \quad (2.4.0)$$

At a certain current density (injected electrons) at the base-side of the collector,  $E(0) = 0$  at the edge of the base-collector junction. This is often referred to as the Kirk threshold current density

$$J_{Kirk} = J_{max} = \frac{2\varepsilon_o\varepsilon_r v_{eff}}{T_c^2} (\phi_{bi} + V_{cb,i}) + qN_c v_{eff} \quad (2.4.0)$$

where  $V_{cb,i}$  is the intrinsic voltage across the base collector junction.  $V_{cb,i} = V_{cb,applied} - I_c \cdot (R_c + R_{ex})$ . At higher  $J_e > J_{max}$  the position of the zero electric field pushes further into the collector. This causes the conduction and valence bands progressively flatten within the collector to a distance  $T_1$ ,  $0 \leq x < T_1$ ,  $E(x) = 0$ , so that the potential barrier for holes is zero. This causes the base to be *pushed out* and hence the base transit time  $\tau_b$  and collector-base capacitance  $C_{cb}$  increase. This is the classical definition of the Kirk effect [18]. As seen from Eqns. ?? ,higher HBT bandwidths are achieved at high current densities -  $J_{max}$  should be increased at a given collector thickness. From Eqn. 2.4.0, this can be done by increasing the  $V_{cb,i}$  and/or increasing the collector doping  $N_c$ . When designing an HBT for use in a digital circuit, two bias conditions need to be considered:  $V_{cb} = 0$ ,  $J_e = J_{max}$  and  $J_e = 0$ . To maximize  $J_e$ , the collector doping should be designed as large as possible while fully depleting the collector at zero bias. From Eqn. 2.4.3 this is satisfied when,

$$N_{c,max} = \frac{2\varepsilon_o\varepsilon_r \phi_{bi}}{qT_c^2} \quad (2.4.0)$$

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and Eqn. 2.4.0 can be rewritten as

$$J_{max} = \frac{4\varepsilon_o\varepsilon_r v_{eff}}{T_c^2} (\phi_{bi} + V_{cb}) \quad (2.4.0)$$

If the transistor is operating at  $J_{kirk}$ , the parasitic voltage drops across the collector and emitter resistances can greatly reduce the intrinsic collector-base voltage.  $J_{kirk}$  can be written as,

$$J_{Kirk} = \frac{4\varepsilon_o\varepsilon_r v_{eff}}{T_c^2} (\phi_{bi} + V_{cb,applied} - I_c(R_c + R_{ex}))$$

$$J_{Kirk} = \frac{4\varepsilon_o\varepsilon_r v_{eff}}{T_c^2} \left( \frac{\phi_{bi} + V_{cb,applied}}{1 + A_E \cdot (R_c + R_{ex}) \frac{4\varepsilon_o\varepsilon_r v_{eff}}{T_c^2}} \right) \quad (2.4.0)$$

At the doping given by Eqn. 2.4.3,  $J_{max}$  is  $2\times$  higher compared to a collector that

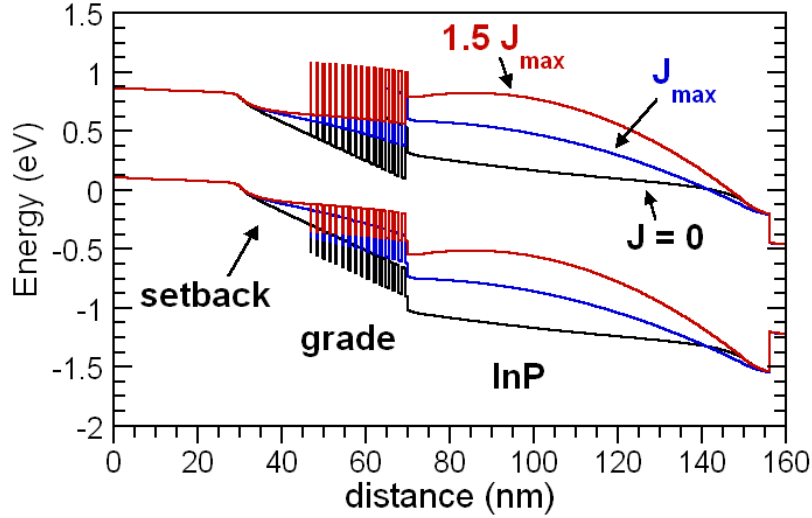


Figure 2.4: Band diagram at  $J=0$ ,  $J_{kirk}$  and  $1.5J_{kirk}$  for  $V_{cb} = 0$

is undoped, greatly influencing logic speed through the use of smaller devices at a given operating current  $I_c$ . A band-diagram for the base-collector junction is shown in Fig. 2.4 for  $J_e = 0$ ,  $J_{max}$ , and  $1.5 J_{max}$  at  $V_{cb} = 0$ . This clearly shows the base pushout regime. Beyond the maximum operating current density, the current gain falls and  $f_\tau$  and  $f_{max}$  decrease.

## 2.5 Device modeling

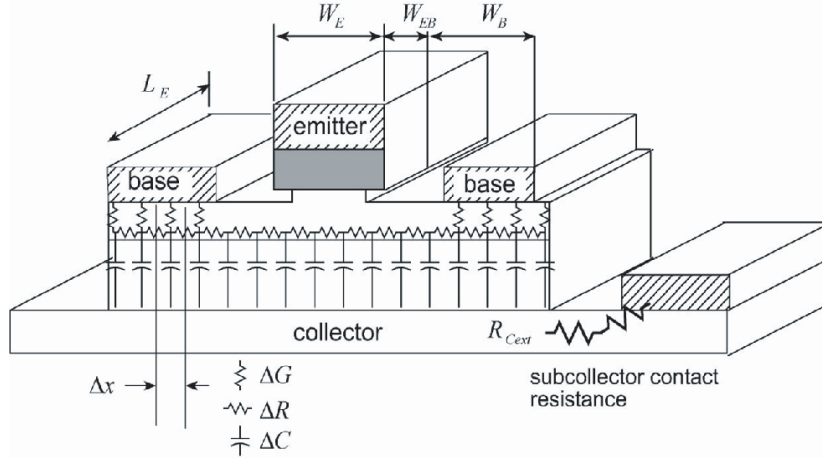


Figure 2.5: Mesa HBT showing distributed device resistances and capacitances

This section describes the small signal modeling schemes used to characterize InP HBTs from UCSB. It makes use of the measured S-parameters of the HBT in order to extract their electron transit time, resistances, and capacitances to create a hybrid- $\pi$  equivalent circuit. It is an accurate, simplified, non-scaleable representation of an HBT compared to the true distributed nature of  $RC$  elements within the device. A SPICE model which is scaleable and physically based and includes bias and frequency dependence is used for circuit design and simulations. Fig. 2.6 shows the T-model equivalent circuit for an HBT. The T-model is physically derived and is intuitive.

$$\alpha(\omega) \approx \alpha_0 \frac{1}{1 + j\omega\tau_b} e^{-j\omega\tau_c} \frac{\sin(2\omega\tau_c)}{2\omega\tau_c} \quad (2.5.0)$$

where  $\alpha_0$  is the DC common base current gain. The T-model while physically based is difficult to work with except for analyzing common base stages. Through a series of transformations detailed in [19], the T-model can be simplified into a hybrid- $\pi$  model which is shown in Fig. 2.7. The hybrid- $\pi$  model shows that the base emitter capacitance consists of the junction capacitance and a diffusion capacitance. This diffusion capacitance,  $C_{diff}$  is an equivalent capacitance derived from the T-model as  $C_{diff} = g_m\tau_f$  where  $\tau_f = \tau_b + \tau_c$  is the total transit time through the base and the collector. The resistance  $R_{be}$  across the base emitter junction is also derived from the T-model and is given as  $R_{be} = \beta/g_m$ .

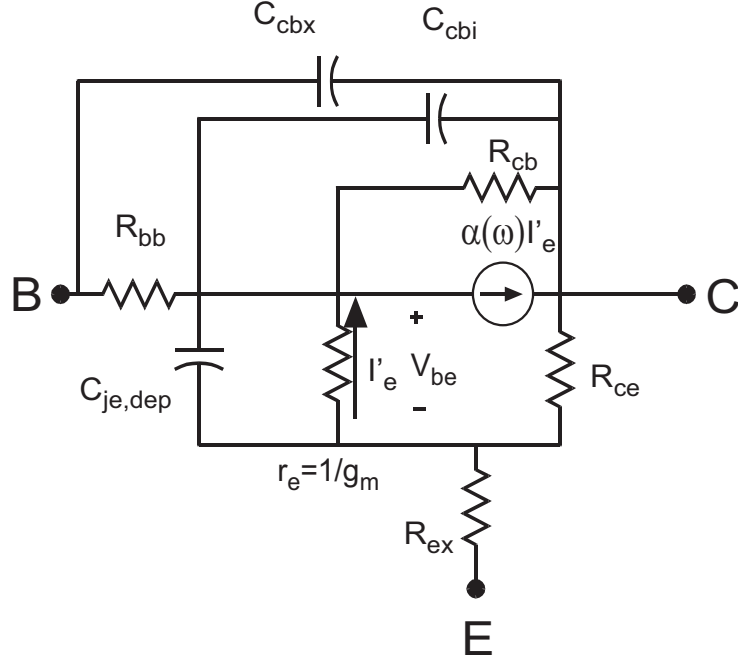


Figure 2.6: T-model equivalent circuit of an HBT

The various capacitances and resistances are extracted from RF (and DC) measurements.

The device transconductance  $g_m$  is given by,

$$g_m(\omega) = \frac{qI_e}{\eta kT} \cdot \exp^{-j\omega\tau_f} \quad (2.5.0)$$

where  $\eta$  is the ideality factor of the collector current, extracted from the gummels as explained below. A non-unity ideality factor can arise when the current-voltage relations deviate from the elementary diode current formulation. This happens due to recombination currents, and thermionic emission over the heterobarrier at the base emitter interface.

The transistor Y parameters are obtained from the 2-port S parameter measurement of the device. At low frequencies,

$$Re(Y_{21})^{-1} = R_{ex} + \frac{R_{bb}}{\beta} + \frac{\eta kT}{qI_e} \quad (2.5.0)$$

$Re(Y_{21})^{-1}$  is plotted for various bias currents.  $\beta = \partial I_c / \partial I_b$  is determined from the low frequency value of  $h_{21}$ . The intercept at  $1/I_e = 0$  gives  $R_{ex} + R_{bb}/\beta$  while  $\eta$

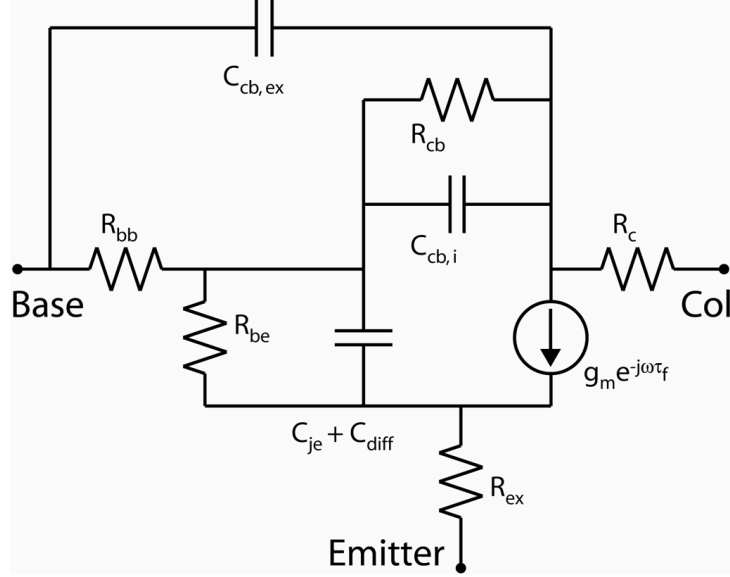


Figure 2.7: Hybrid-pi equivalent circuit HBT model

can be determined from the slope. The collector-base conductance  $1/R_{cb}$  and total collector-base capacitance  $C_{cb}$  are determined from the real and imaginary parts of  $Y_{12}$  at low frequencies,

$$Y_{12} = \left( \frac{1}{R_{cb}} + \omega^2(C_{be} + C_{cb,i})C_{cb,i}R_{bb} \right) + j\omega(C_{cb,i} + C_{cb,ex}) \quad (2.5.0)$$

This method involves RF measurements of transistors and is accurate but time consuming. An extraction from DC measurements is developed in this thesis. This involves extraction from Gummel plots, which are logarithmic plots of  $I_C$  and  $I_B$  vs.  $V_{BE}$ . The collector current can be expressed as,

$$I_C = I_{CO} e^{\frac{q}{kT_A} [V_{BE,i} - (T - T_A) \frac{\partial V_{BE,i}}{\partial T}]} \quad (2.5.0)$$

where  $T_A$  is the ambient temperature,  $\sim 300\text{K}$  and  $V_{BE,i}$  is the intrinsic base emitter voltage given by

$$V_{BE,i} = V_{BE} - I_C \cdot (R_{ex} + R_{bb}/\beta) \quad (2.5.0)$$

where  $V_{BE}$  is always the measured base emitter voltage. The effect of the emitter and base resistances are seen as deviations from the exponential dependence in the gummel plot (see Fig. ??). In a linear plot, the resistances cause a leanover in the

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current in the diode I-V characteristics. The change in  $V_{BE}$  as a result of temperature change in the device is given by,

$$\delta V_{BE} = -\Theta_{th} \cdot \frac{\partial V_{BE}}{\partial T} \cdot I_C \cdot \delta V_{CE} \quad (2.5.0)$$

where,  $\Theta_{th}$  is the thermal resistance usually expressed in  $^{\circ}\text{K}/\text{mW}$ . The effect of

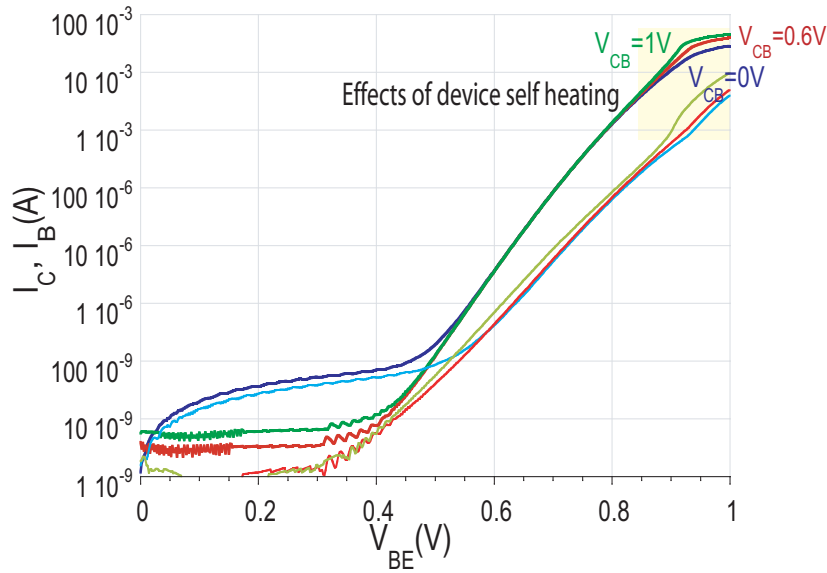


Figure 2.8: Effect of device heating, with  $V_{CB}$ , in the gummel plot

temperature on  $V_{be}$  is particularly large at high current densities and can be seen in the as measured gummel curves in Fig. 2.8. The device heating is due to increasing reverse collector base biases and therefore  $I_C$  in the gummels is dependent on the value of  $V_{CB}$ . In a well designed DHBT, where the early voltages are very high ( $\geq 1500\text{V}$ ),  $I_C$  should not have any  $V_{CB}$  dependence in the active region of operation (the effect of collector resistance is ignored and will be revisited in Chapter ?? ). In order to accurately extract  $R_{ex}$  from the Gummels, this thermal dependence of  $V_{BE}$  has to be removed, and it is then named as  $V_{BE}^{isothermal}$ .  $I_C$  vs  $V_{BE}^{isothermal}$  should therefore have no dependence on  $V_{CB}$ . A method(unpublished) to extract isothermal gummels is suggested.

From Eqn. 2.5, the isothermal base emitter voltage is written as,

$$V_{BE}^{isothermal} = V_{BE} - \Theta_{th} \cdot \frac{\partial V_{BE}}{\partial T} \cdot I_C \cdot \delta V_{CE} \quad (2.5.0)$$

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The detailed derivation is skipped but  $V_{BE}^{isothermal}$  can be expressed in terms of  $V_{CB}$  and  $V_{BE}$  as,

$$V_{BE}^{isothermal} = \frac{V_{BE}}{1 + \Delta V_{BE}/\Delta V_{CB}} - \frac{V_{CB}}{1 + \Delta V_{CB}/\Delta V_{BE}} \quad (2.5.0)$$

This suggests that  $\Delta V_{BE}/\Delta V_{CB}$  should be calculated at each  $I_C$ . However, in the Gummels measurement procedure,  $I_C$  is measured at various applied base emitter voltages  $V_{BE}$ . An approximate value of  $\Delta V_{BE}/\Delta V_{CB}$  could be used for different ranges of  $I_C$  but due to this quantization error, kinks are seen in the resulting isothermal gummel curves. A simple mathematical analysis is shown below.

$$\begin{aligned} I_C &\sim I_C(V_{BE}, V_{CB}) \\ \partial I_C &= 0 \text{ since} \\ \frac{dV_{BE}}{dV_{CB}} &\text{ is needed at constant } I_C. \text{ This leads to} \\ \frac{dV_{BE}}{dV_{CB}} &= - \frac{\left. \frac{\partial I_C}{\partial V_{CB}} \right|_{V_{BE}}}{\left. \frac{\partial I_C}{\partial V_{BE}} \right|_{V_{CB}}} \end{aligned} \quad (2.5-3)$$

This approximates to a linear interpolation for small, discrete values of  $\Delta V_{BE}$  and  $\Delta V_{CB}$ .  $V_{BE}^{isothermal}$  is thus calculated by using small values of  $\Delta V_{BE}$  and  $\Delta V_{CB}$ . Fig. 2.9 shows the isothermal gummels and as expected,  $I_C$  does not depend on  $V_{CB}$ . From the isothermal plot,  $R_{ex} + R_{bb}/\beta$  is extracted as shown in Fig. 2.9.  $\beta$  is the current gain of the device known from the gummel curves. Usually  $\beta$  is large enough that the contribution of  $R_{bb}/\beta$  is neglected. The resistance seen in the emitter is thus plotted in Fig. 2.10.

Furthermore this method is also an extraction procedure for the thermal resistance of the device,  $\Theta_{th}$ . which is seen from Eqn. 2.5 to be

$$\Theta_{th} = - \frac{\delta V_{BE}}{\frac{\partial V_{BE}}{\partial T} \cdot I_C \cdot \delta V_{CE}} \quad (2.5-3)$$

where  $\phi = \frac{\partial V_{BE}}{\partial T}$  is called the thermal-electrical feedback coefficient, expressed in  $mV/^\circ K$ . It is a property of the transistor and is approximately a constant ( $\approx 2mV/^\circ K$  for InPDHBTs).

The total delay  $\tau_{ec}$ , that electrons experience through the HBT is,

$$\frac{1}{2\pi f_\tau} = \tau_c + \tau_b + C_{cb} \cdot (R_{ex} + R_c) + \frac{\eta k T}{q I_e} (C_{cb} + C_{je}) \quad (2.5-3)$$



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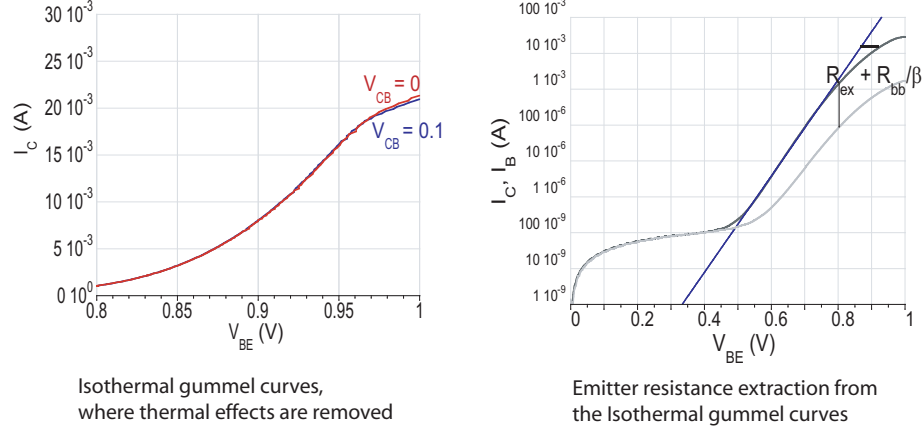


Figure 2.9: Isothermal gummel plot and effect of emitter and base resistances

$f_\tau$  is extracted from  $h_{21}$  at different bias currents and  $C_{cb}$ ,  $R_{ex}$ , and  $g_m$  have already been determined.  $\frac{1}{2\pi f_\tau}$  is plotted for the various bias currents at which  $f_\tau$  is measured. The slope gives  $C_{je}$  since  $C_{cb}$  is already obtained from 2.5.  $R_{ex}$  is known from 2.5. with knowledge of  $R_c$ ,  $\tau_f = \tau_c + \tau_b$  and hence  $C_{diff}$  can be determined from the  $1/I_e = 0$  intercept.  $R_c$  is usually calculated for the HBT geometry knowing the values, from TLM measurements, of contact resistivity  $\rho_c$  and sheet resistance  $R_{sh}$ . This not accurate especially when  $R_c$  is dominated by other than the contact resistance. An alternate method is explored in this thesis using the DC  $I_C$ -  $V_{CE}$  characteristics. From the Ebers-Moll model for a bipolar transistor,  $V_{CE}$  is given by

$$V_{CE} = \frac{kT}{q} \ln \frac{1 + \frac{I_C}{I_B} \cdot (1 - \alpha_r)}{\alpha_r (1 - \frac{I_C}{I_B \cdot \beta_F})} + I_C (R_{ex} + R_c) + I_B \cdot R_{ex} \quad (2.5.-3)$$

where  $\beta_F$  is the forward current gain and  $\alpha_r = \frac{\beta_r}{\beta_r + 1}$  as defined in the Ebers-Moll equations [20]. If  $\beta = I_C/I_B$  is kept constant for various values of  $I_C$ , term 1 in Eqn. 2.5 is constant. From the  $I_C$ - $V_{CE}$  characteristics,  $V_{CE}$  is plotted vs.  $I_C$ , keeping  $\beta$  a constant. The slope of this curve gives  $R_c + R_{ex} + R_{ex}/\beta$ .  $R_{ex}$  has already been determined from  $\text{Re}(Y_{11})$  and from the gummel extraction procedure described.  $R_c$  can now be extracted. The variation of  $V_{CE}$  with current is plotted for several values of  $\beta$ . This method extracts  $R_c$  from the saturation regime of transistor operation where  $I_C$  increases with  $V_{CE}$ . This is far from the usual operation of the transistor. The base resistance is determined by comparing the real part of  $Y_{11}$  of the measured

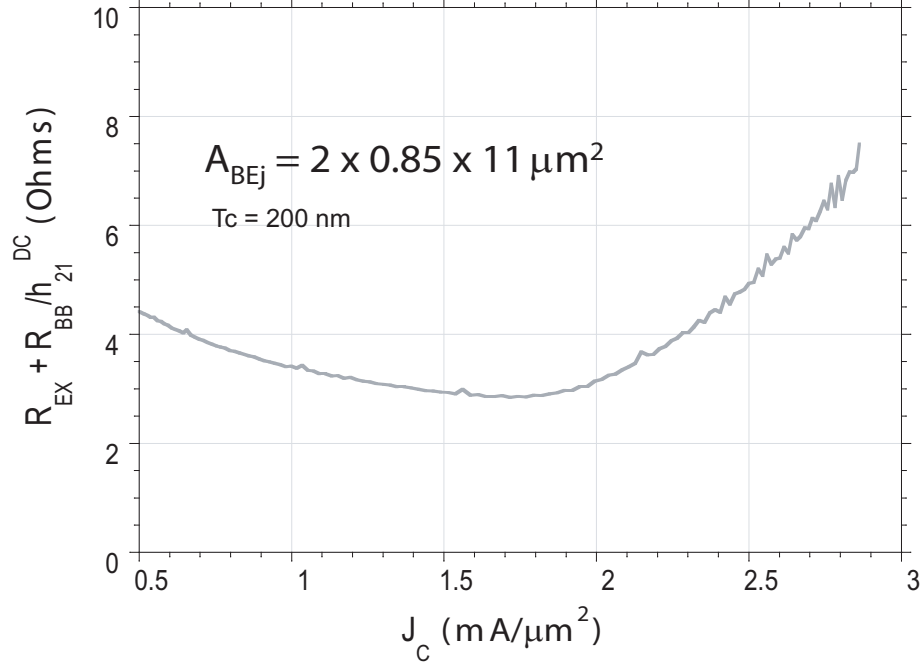


Figure 2.10: Extracted emitter and base resistance

HBT data to the equivalent circuit from the following relation,

$$Re(Y_{11}) \approx \frac{1}{R_{be}} + \omega^2(C_{je} + C_{diff})^2 \cdot R_{bb} \quad (2.5.-3)$$

Through the hybrid- $\pi$  model,  $R_{bb}$  is adjusted to make the quadratic frequency dependence of  $Re(Y_{11})$  match the measured trend. Once determined,  $R_{bb}/\beta$  is known.  $C_{cb,i}$  is similarly determined from  $ReY_{12}$  in Eqn. 2.5. Mason's unilateral gain  $U$  is only influenced by  $C_{cb,i}$ . By simultaneously monitoring  $Re(Y_{12})$  and  $U$ ,  $C_{cb,i}$  is adjusted while keeping the total  $C_{cb}$  constant to match the measured data.

Analysis of the hybrid- $\pi$  network gives  $Im(Y_{12}) = \omega C_{cb}$ . From the S-parameters measured,  $Y_{12}$  can be extracted. This is used to extract the total  $C_{cb}$ . From the C-V

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data at  $I_c = 0$  mA, the doping density can be derived as follows.

$$C_{cb} = -\frac{Q_c}{V_{bc}} \quad (2.5.-2)$$

$$dQ_c = qN(x_d)dx$$

In general, the doping density can be expressed as,

$$N(x_d) = -\frac{Cdx}{qdV} = -\frac{C}{q} \frac{dC}{dV} \frac{dx}{dC} \quad (2.5.-3)$$

$$N(x_d) = \frac{dC_{cb}/dV_{cb}}{q\epsilon A_c^2 C_{cb}^3} = -\frac{2}{q\epsilon A_c^2} \cdot \frac{dV}{d(1/C_{cb}^2)}$$

The depletion region edge is given by

$$x_d = \frac{\epsilon A_c}{C_{cb}} \quad (2.5.-4)$$

. Therefore, the doping density can be plotted as a function of depletion distance in the collector. This neglects the effect of Deby length on the doping profile extracted [21]. Fig. 2.11 shows on a Smith chart measured HBT S-parameters and those of its equivalent circuit. The  $f_\tau$  and  $f_{max}$  of the hybrid- $\pi$  equivalent circuit is consistent with the value extrapolated for the HBT.

## 2.6 HBT delays within digital ICs

The HBT figures-of-merit  $f_\tau$  and  $f_{max}$  describe the maximum bandwidth for a single device. They are of limited value in predicting digital logic speed. A standard benchmark for digital logic figure-of-merit, for a device technology, is the bandwidth of a static frequency divider shown in Fig. 2.12. It is a master-slave (M-S) flip-flop consisting of two series connected latches that are clocked out of phase  $180^\circ$  so that the input is transparent at the output only at the falling edge of the clock. As shown in the timing diagram in Fig. 2.12, when  $\bar{Q}$  is connected to D, the output  $= f_{CLK}/2$ . MS latches are utilized as retiming elements for data synchronization and their maximum toggle rate often limits circuit bandwidth. For this reason, static dividers are a more realistic benchmark circuit in comparison to the narrow-band operation of dynamic frequency dividers and ring oscillators.

The propagation delay through the latch is dependent upon the combined charging times of the capacitances in the signal path. By modeling the latch as an n-port linear network having no inductors, the method of open circuit time constants (MOTC) can be used to evaluate the time constants associated with the poles  $a_n$  of

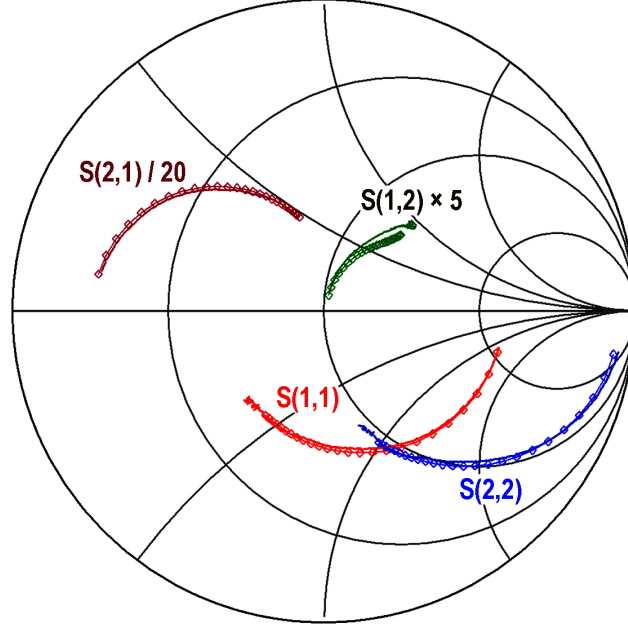


Figure 2.11: Measured (solid line) and simulated S-parameters (data points) of the HBT and the extracted hybrid- $\pi$  equivalent circuit

the system. The transfer function for such a network is given by,

$$\frac{V_{out}}{V_{in}} = A_{v,midband} \left( \frac{1 + b_1s + b_2s^2 + \dots + b_ns^n}{1 + a_1s + a_2s^2 + \dots + a_ns^n} \right) \quad (2.6.0)$$

In order to utilize MOTC, the passive and active components of the network must behave linearly. Each arm of the differential pair (say  $Q_1$  and  $Q_2$ ) in Fig. 2.13, conduct current in alternate clock cycles. Thus the small signal values of  $g_m$  and  $C_{je}$  are no longer valid. To satisfy this requirement, the HBT transconductance  $g_m$  and diffusion capacitance  $C_{diff}$  of those devices operating as part of a differential pair (where the voltage swing across the base emitter junction is  $\approx \Delta V_{logic} \gg kT/q$ ) are modified,

$$G_{m, large-signal} = \frac{\Delta I_o}{\Delta V_{be}} \approx \frac{1}{R_{Load}} \quad (2.6.1)$$

$$C_{diff, large-signal} = G_m \tau_f \quad (2.6.2)$$

$$C_{je, large-signal} = \frac{1}{V_1 - V_2} \int_{V_1}^{V_2} C_{je}(V) dV \quad (2.6.3)$$

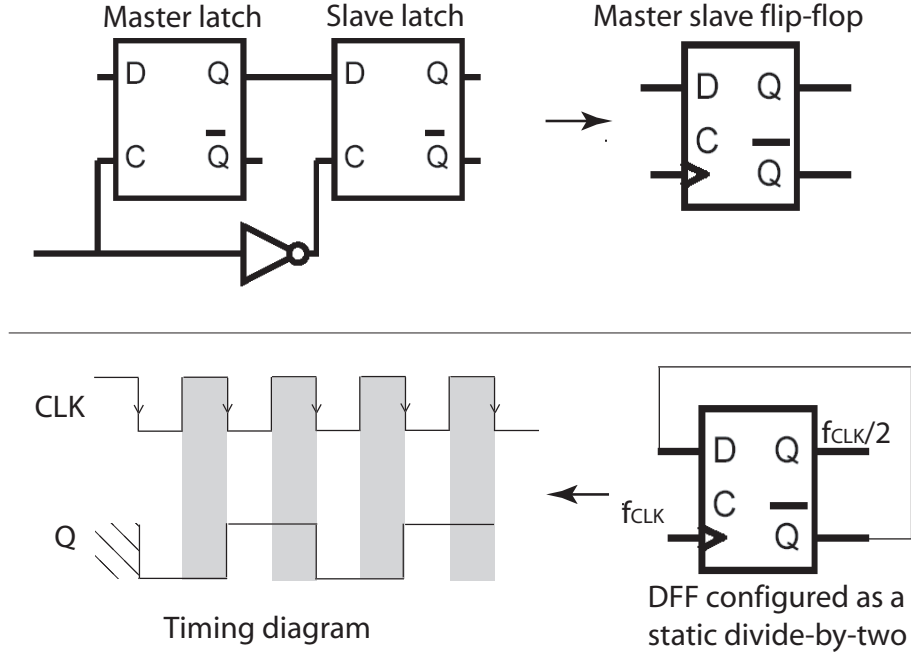


Figure 2.12: Static frequency divider and its timing diagram

Large signal operation greatly reduce  $G_m$  and  $C_{diff}$  by  $\sim 1:10$ . The depletion capacitance  $C_{cb}$ , is not altered and the charging (and discharging) of this capacitance constitutes a major delay in a digital circuit. From the method of time constants [?],  $a_1$  is,

$$a_1 = R_{11}^0 C_1 + R_{22}^0 C_2 + R_{33}^0 C_3 + \dots R_{nn}^0 C_n \quad (2.6.3)$$

where  $R_{nn}^0$  is the effective resistance across the terminals of  $C_{nn}$  with other capacitances treated as open (for example, see [22]). The propagation delay is typically defined as the time required to effect a change in the output node when the input is toggled. Assuming  $(1 - \exp(-t/\tau))$  charging behavior,  $T_{prop} = a_1 \ln(2)$ . However, to the level of accuracy of the assumptions used in the analysis,  $T_{prop} = a_1/2$ . Fig. 2.13 shows the significant resistive and capacitive delay elements in the signal

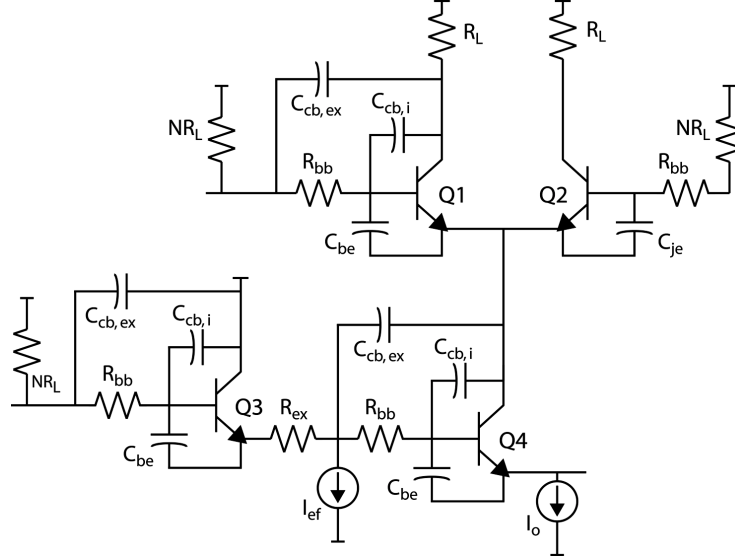


Figure 2.13: Delay path of a CML static frequency divider during clock transition

path of a two level CML flip-flop. The propagation delay is simplified to

$$\begin{aligned}
 2T_{prop} &\cong N(\Delta V_{logic}/I_o)C_{cb3} + R_{bb}C_{cbi3} \\
 &+ [2(R_{ex3} + 1/g_{m3}) + (\Delta V_{logic}/I_o)]C_{cb4} + 2C_{cbi4}R_{bb4} \\
 &+ (C_{je4} + I_o\tau_f/\Delta V_{logic})(R_{bb4} + R_{ex3} + 1/g_{m3}) \\
 &+ [C_{je2} + C_{je1} + (I_o\tau_f/\Delta V_{logic})](\Delta V_{logic}/I_o) \\
 &+ (N + 1)(\Delta V_{logic}/I_o)C_{cb1} + R_{bb1}C_{cbi1}
 \end{aligned} \tag{2.6.0}$$

where the devices are operating in the following modes:  $Q_3$  emitter-follower,  $Q_4$  common-emitter,  $Q_1$  common-base, and  $Q_2$  cut-off.  $N$  denotes the fan-out of identically connected gates in a larger circuit. As seen from Eqn. 2.6.0, the base and collector transit times play a relatively minor role in logic speed, in comparison to their strong contributions to  $f_\tau$ . The most significant delays in the latch are from charging the depletion capacitances over the logic swing,  $(C_{je} + C_{cb})\Delta V_{logic}/I_o$ , where  $\Delta V_{logic}/I_o = R_L$ . This can be expressed in terms of the current density and transistor sizes as

$$(C_{je} + C_{cb})\frac{\Delta V_{logic}}{J_e \cdot A_e} = \hat{C}_{je} \cdot \frac{\Delta V_{logic}}{J_e} + \hat{C}_{je} \cdot \frac{A_C}{A_E} \cdot \frac{\Delta V_{logic}}{J_e} \tag{2.6.0}$$

To minimize these capacitive delays, small devices should be used operating a current density  $J_e$  close to  $J_{Kirk}$  and the collector to the emitter area ration( $A_C/A_E$ )

## CHAPTER 2. INP DHBT THEORY AND DESIGN

must be made small which is the objective of this dissertation. From the above expression 3.3,  $C_{cb}/I_c$  expressed in ps/V is an important metric to be considered while comparing digital circuits in various technologies.

$\Delta V_{logic}$  is dependent on the input noise margin which is influenced by  $R_{ex}$ . It is not evident from the gate delay expression, but  $R_{ex}$  has a large indirect effect on the maximum toggle rate. For the differential pair in Fig. 2.14, the differential switching current is described by,

$$I_{c1} - I_{c2} = I_o \tanh \left[ \frac{q(V_1 - V_2)}{2kT} - \frac{I_{e1} - I_{e2}}{2} \cdot R_{ex} \right] \quad (2.6.0)$$

With  $R_{ex} \sim 0$ , the current is completely switched when  $V_1 - V_2 = 6kT/q$ . As  $R_{ex}$  increases, at a fixed current  $I_o$ , increasing input voltage is needed to completely switch the differential pair. When the voltage drop across  $R_{ex}$  is  $6kT/q$ , more than  $10kT/q$  potential difference is needed to switch the currents. If we assume an HBT junction temperature rise of  $\sim 60^\circ\text{C}$  when these devices operate in a larger circuit, the potential difference required to switch only the base-emitter junction is  $\sim 200$  mV. Typically,  $V_1 - V_2 = \Delta V_{logic} = 300$  mV for ECL and CML based latches. Thus the digital noise margin is  $\sim 100$  mV  $\approx 4kT/q$ .

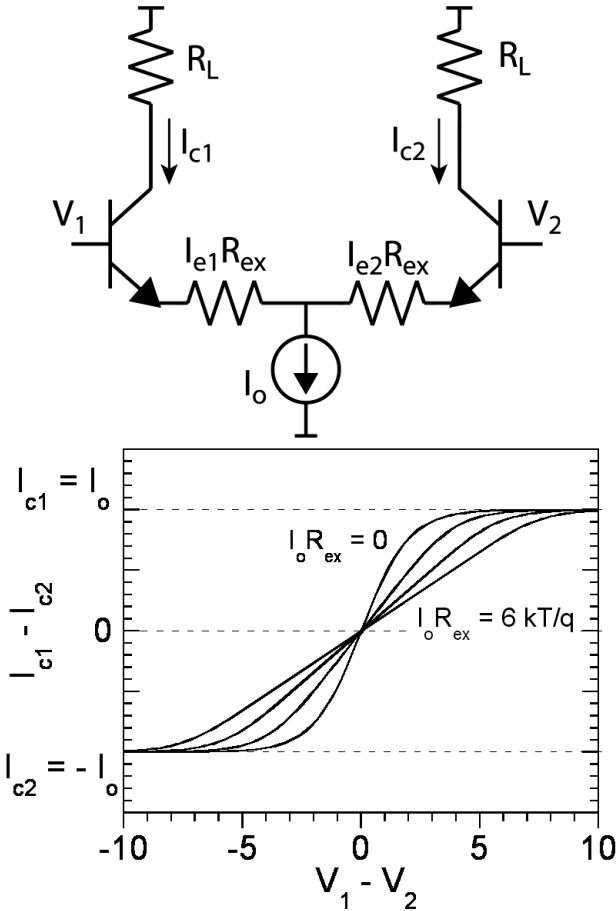


Figure 2.14: Current flow of a differential pair in the presence of emitter resistance –  $I_o R_{ex} = 0, 2kT/q, 4kT/q,$  and  $6kT/q$ .  $V_1 - V_2$  is normalized to  $kT/q$ .



## 2.7 HBT scaling principles

The previous sections reviewed the relevant HBT transit and  $RC$  delays associated with discrete device performance and those of a digital latch. In order to improve HBT as well as analog and digital IC speed, all significant capacitances and transit times must be simultaneously reduced. This is summarised as the scaling laws for a HBT [3].

As seen in the earlier sections,  $f_\tau$  is dominated by  $\tau_f$  while increasing current density decreases digital delay. If  $T_c$  and  $T_b$  are decreased by  $\gamma$  and maintaining all device dimensions constant,

$$\begin{aligned} \tau_c &\downarrow \gamma^2 \\ \tau_b &\downarrow \gamma^2 \\ J_{kirk} &\uparrow \gamma^2 \\ C_{cb} &\uparrow \gamma \\ R_{bb} &\uparrow \approx \gamma \end{aligned} \tag{2.7.-3}$$

If the collector junction areas are also reduced by  $1:\gamma^2$ ,  $C_{cb}$  will also decrease by  $\gamma$ . To reduce  $R_{bb}$ , the base contact resistivity would have to be decreased by  $\sim \gamma^2$ . If the base ohmic contacts lie above the collector-base junction, their width must be reduced  $1:\gamma$  to obtain the requisite reduction in  $C_{cb}$ ; this necessitates a further reduction in the base contact resistivity  $\rho_{c,b}$ . However, this is very difficult and limited by the doping in the base, the work function of the contact metal used and by surface preparation prior to metal deposition. Therefore, one of the biggest challenges to vertical device scaling is the scaling of the collector base capacitance. Independent scaling of the emitter and collector junction is necessary and is the main subject of this dissertation.

It is also seen from Eqn. 2.7.-3 that current density increases with vertical scaling. If the emitter junction dimensions are kept constant  $I_e$  increases. This increases the power consumption in the device and thermal effects assume significance. Therefore it is equally important to scale the emitter junction area. The power consumption in the divider is very approximately,

$$P_L \propto \Delta V_{logic} \times J_e \cdot A_{je} \approx \left(6 \frac{kT}{q} + 2J_e A_{je} R_{ex}\right) \times J_e L_e W_{je} \tag{2.7.-3}$$

The width of the junction area is limited by the lithographic and process tolerances. To decrease power consumption by  $1:\gamma$  while maintaining high high current density, means decreasing the emitter (and device) length, so that the emitter junction area  $A_{je} \downarrow \gamma^2$ . This is especially important for low power logic. As seen in the previous

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section, a significant component of the delay in logic circuit is

$$\hat{C}_{cb} \cdot \frac{W_{bm}}{W_{je}} \cdot \frac{\Delta V_{logic}}{J_e} + C_{cb,pad} \cdot \frac{\Delta V_{logic}}{J_e L_{je} W_{je}} \quad (2.7.-3)$$

where  $\hat{C}_{cb}$  is the device capacitance per unit area (not including  $C_{cb,pad}$ , the access pad capacitance). A certain access pad area ( $\approx 2 \mu\text{m}^2$ ) is needed for the device to reliably contact the metal interconnect and to maintain a low metal access resistance. This base pad cannot be scaled. As seen from Eqn. 3.3, as the length is decreased,  $C_{cb,pad}$  becomes a significant fraction of the total  $C_{cb}$ . Chapter ?? addresses this through the use of implanted subcollectors.

In a standard triple-mesa DHBT process, decrease in the junction area is accompanied by an increase in the emitter resistance, the emitter contact area has decreased as well. Besides affecting logic speed and  $f_\tau$ , as seen in the precious section an increase in  $R_{ex}$  severely impacts the noise margin of the logic circuit. If  $\rho_{ex}/A_{em} \cdot J_e A_{ej}$  is to remain a constant where  $A_{em}$  is the area of the emitter contact,  $\rho_{ex}$  must decrease in proportion to increase in  $J_e$ . This is a process challenge and addressed by the use of highly doped InAs layers[?]. An alternate method to maintain  $R_{ex}$  constant, is to decrease  $A_{je}$  and maintain  $A_{em}$  constant. This can be done by severe undercutting of the emitter base junction which presents process difficulties or by emitter regrowth discussed in Chapter ?? .

Vertically scaling the collector thickness decreases the breakdown voltages,  $BV_{CEO}$  and  $BV_{CBO}$ . The breakdown voltage is limited by several factors in a DHBT. Firstly the term 'breakdown voltage' needs clarification. Traditionally, open base or open emitter voltage at a certain current level (  $50 \mu\text{A}$ ) are often reported. At these current levels, leakage current due to passivation or surface states limit the maximum voltage attained. If the setback and grade layers are thick, impact ionisation could occur in these regions and thus the advantage of having wideband gap InP collector is lost. (It is unclear at this time whether breakdown is due to impact ionisation or tunneling.) Since operating  $J_e$  increases with vertical scaling, the maximum reliable power density associated with a *safe operating area*  $P/A_e \sim J_e V_{ce} \propto \gamma^2 V_{ce}$  is a more significant applied voltage limit than the low-current breakdown voltages  $BV_{CEO}$  or  $BV_{CBO}$ .

The total emitter-base capacitance  $C_{je}$  is given by,

$$C_{je} \approx \frac{\kappa_1 L_e W_e}{T_{eb}} \quad (2.7.-3)$$

If the emitter junction is scaled laterally, to first order  $C_{je}$  is reduced  $\gamma^2 : 1$  (where  $\kappa_1$  is a constant). Scaling requirements for the emitter depletion thickness  $T_{eb}$  are not easily summarized here and detailed analysis for scaling  $T_{eb}$  is reported in [3].

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Table 2.1: Summary of simultaneous parameter scaling for a  $\gamma:1$  increase in HBT and circuit bandwidth

key device parameter	required change
collector depletion layer thickness	decrease $\gamma:1$
base thickness	decrease $\sqrt{\gamma}:1$
emitter-base junction width	decrease $\gamma^2:1$
collector-base junction width	decrease $\gamma^2:1$
emitter depletion thickness	decrease $\gamma^{1/2}:1$
emitter contact resistivity, $\rho_{ex}$	decrease $\gamma^2:1$
emitter current density	increase $\gamma^2:1$
base contact resistivity – if contacts lie above B-C junction	decrease $\sim \gamma^2:1$
base contact resistivity – if contacts do not lie above B-C junction	unchanged
bias currents and voltages	unchanged

Lastly, reductions to device thermal resistance need to be considered. For discrete HBTs, a significant fraction of the heat generated in the collector is removed through the base and into the emitter metal. In large integrated circuits, the heat-sinking through the emitter is much less effective.. The thermal resistance normalized to the emitter junction area  $\theta_{JA}A_{je}$  must be reduced in proportion to the square of the circuit bandwidth  $\gamma^2 : 1$ . Thinning the  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  subcollector layer is imperative [14] to decreasing the device thermal resistance.

What is the impact of this scaling on ECL (and CML) logic speed? With a  $\gamma:1$  scaling, the collector thickness  $T_c$  is reduced  $\gamma:1$ , the current density increased  $\gamma^2:1$ , and the dominant delay  $C_{cb}\Delta V_{logic}/I_c$  reduced  $1:\gamma$ , **only if** the access pad capacitance  $C_{cb,pad}$  is eliminated. The parasitic voltage drop  $R_{ex}I_c = \rho_{ex}J_e$  remains constant if  $\rho_{ex}$  is reduced rapidly by  $1:\gamma^2$ . The scaling laws for a  $\gamma$ -fold increase in bandwidth are summarised in Table 2.1

## 2.8 HBT scaling limits, and solutions explored in this work

In order to increase circuit bandwidth, the transit times and capacitances of the device must be reduced while maintaining constant (total) resistances, currents, and  $g_m$ . This is realized by thinning the base and collector layers, narrower emitter and collector junctions, increased operating current density, and reduced contact  $\rho_c$  and

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sheet  $\rho_s$  resistivities.

Emitter contacts as narrow as  $0.4 \mu\text{m}$  can be achieved through standard optical lithography and evaporated metal-lift-off techniques. If the emitter contact is to be further scaled, advanced photoresist processes are needed to resolve smaller features using i-line stepper lithography. The height of the contact will be limited by the aspect ratio attainable from the e-beam metal evaporator. An electroplated emitter contact, or one formed through metal-sputtering and dry-etch, are alternative ways of producing tall, straight, narrower features. For these narrower contacts, the undercut of the emitter mesa becomes increasingly difficult to control. Dielectric sidewall spacer processes allow for very thin emitter semiconductor layers, minimizing the undercut during mesa formation. Furthermore, the spacer eliminates the need for evaporated self-aligned base contacts, where instead electroplating or metal-sputter dry-etch processes are used. These advanced processes are under development and are discussed in [23, 24]. As discussed, narrow emitter contacts lead to higher emitter resistance. Emitter contact resistivity, to  $N^+$   $\text{In}_{0.85}\text{Ga}_{0.15}\text{As}$ , as low as  $10 \Omega \cdot \mu\text{m}^2$  is now standard at UCSB [13]. It is not clear how  $\rho_c$  can be further reduced by conventional deposition techniques. Advanced materials engineering such as the use of in-situ MBE grown ErAs ohmics may be beneficial. Regrown emitter junction HBTs have also been developed at UCSB as an alternative, where the ohmic contact is much larger than the emitter junction for reduced access resistance.

Thinner base and collector epitaxial layers can be achieved through growth. The base sheet resistance is  $\propto T_b^{-1}$ , and to keep it unchanged the bulk resistivity ( $\rho_{bulk}$ ,  $\Omega \cdot \text{cm}$ ) must be decreased through increased through base doping. As the doping approaches  $7 \cdot 10^{19} \text{cm}^{-3}$ , the hole mobility decreases and the doping-mobility product remains relatively constant. This is a challenge that can be addressed through the use of extrinsic bases described in brief in Chapter ?? The InP DHBTs reported here utilize a 30 nm base and doping grade from  $7 - 4 \cdot 10^{19} \text{cm}^{-3}$ . Through the appropriate choice of metal workfunction and highly doped semiconductor the contact resistivities are decreased. The Fermi level at the semiconductor surface is pinned due to a combination of surface states, native oxides, and metal-semiconductor diffusion at the interface. Surface preparation techniques to minimize their presence have been developed, and low values of contact resistance  $\rho_c \approx 10 \Omega \cdot \mu\text{m}^2$  values have been achieved. The resulting transfer length of the base contact is  $\sim 130 \text{nm}$ . Narrower collector junctions require narrower base contacts. A minimum width  $\sim L_t$ , the ohmic transfer length, should be maintained to prevent exponential increases to the contact resistance  $R_{b,cont}$ . These thin, narrow contacts have high access resistance and inductance that influence HBT performance. Furthermore circuits employing these narrow contacts are very difficult to yield in a manufacturing environment.

Therefore in order to satisfy the scaling requisites for the collector junction, im-

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planted collector processes have been developed and is the main topic of this dissertation. The non scaleable collector base access pad capacitance is identified as a major parasitic. This parasitic capacitance is eliminated through the use of implanted subcollectors. Independent scaling of the collector is highly desirable, within a simple, manufacturable process. A fully implanted pedestal and subcollector transistor is demonstrated where the extrinsic collector base capacitance is greatly reduced.

### 2.9 The scaled HBT

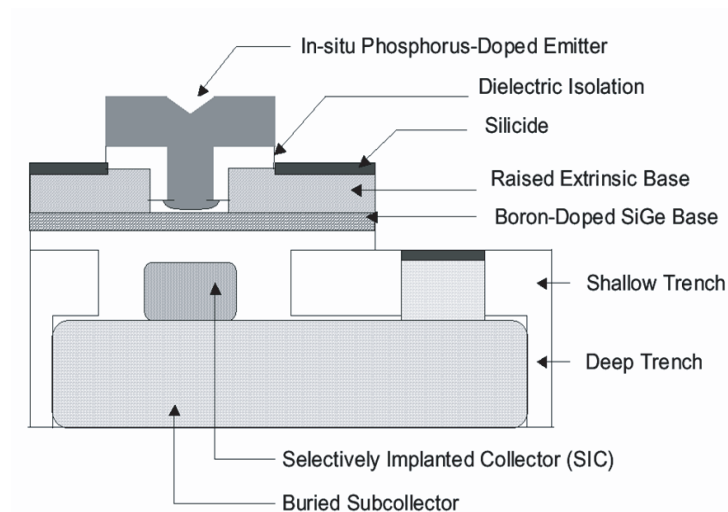


Figure 2.15: Modern SiGe HBT

Compared to modern SiGe HBT, InP HBT manufacturing is primitive. The SiGe HBT shown in Fig. 2.15, has buried subcollectors for zero pad capacitance,  $N^+$  pedestal for lateral collector scaling, deep dielectric trenches for device isolation, thick extrinsic base and buried base ohmics for reduced  $R_{bb}$ , and submicron regrown emitters for emitter-base junction scaling while simultaneously maintaining wide emitter contacts. It has a planar geometry and large scale ICs are possible [?]. Due to this extreme scaling and reduction in relevant parasitics, SiGe HBTs are as fast as their InP counterparts for digital logic applications. Static frequency dividers of  $\sim 100$  GHz have been demonstrated with devices having cut-off frequencies of  $\sim 300$  GHz. This serves as the motivation for scaling InP HBTs and in this thesis, technologies are developed for InP HBTs, to parallel the SiGe HBT manufacturing process.

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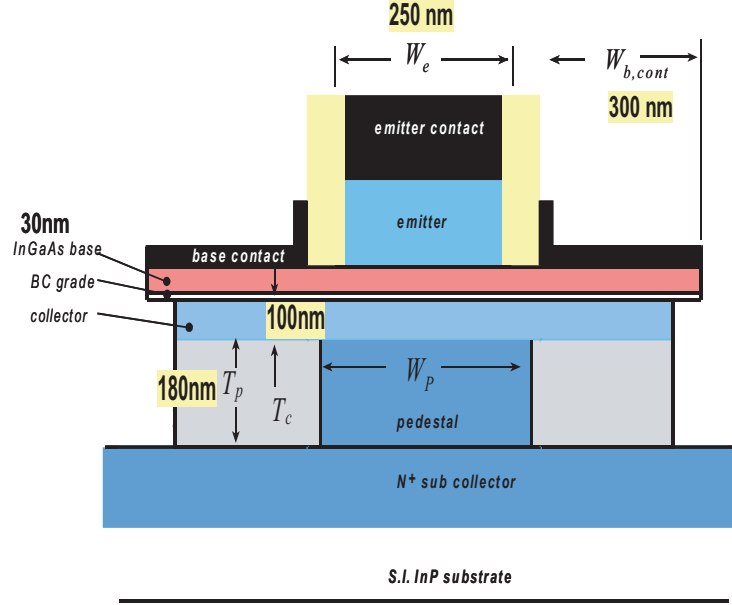


Figure 2.16: Scaled mesa HBT with collector pedestal

The bulk of this thesis examines technologies to reduce the base collector capacitance. In this sub-section, the expected performance enhancement with reduction in  $C_{cb}$  is calculated for highly scaled HBTs. A scaled mesa HBT is shown in Fig. 2.16 employing an emitter sidewall [23] and implanted pedestal-subcollector technology developed in this dissertation. Table 2.2 compares standard figures of merit for a device with 250nm emitter and 300nm base, with and without the implanted collectors.

This calculation assumes all contact resistivities  $\rho_c \sim 10 \Omega \cdot \mu\text{m}^2$ , current density  $J_{kirk}$  is  $13 \text{ mA}/\mu\text{m}^2$ , wiring delay of 0.3 ps, and the width of the emitter junction

Table 2.2: Expected performance of a scaled InP DHBT with and without implanted collectors

Standard Figures of merit	standard mesa DHBT	mesa DHBT with implanted collectors
$f_\tau$	490 GHz	550 GHz
$f_{max}$	600 GHz	800 GHz
$f_{clock}$	180 GHz	270 GHz
$C_{cb}/I_c$	0.4 ps	0.2 ps

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$W_{je} = 0.25 \mu\text{m}$ . The other vertical and lateral dimensions are as in Fig. 2.16. All other parameters are drawn from the state of the art UCSB mesa process [13]. Assuming that the breakdown mechanism is impact ionisation or tunneling, the breakdown voltages ( $\approx 4V$ ) are not expected to be different for the two technologies. The power consumption of digital circuits at a given bandwidth, is expected to be much lower for the mesa DHBT with implanted collectors.

It is seen from Table 2.2 that reduction in  $C_{cb}$  tremendously improves  $f_{max}$  and the maximum digital circuit speed  $f_{clock}$ , while the difference to  $f_{\tau}$  is marginal. As discussed in the prior sections,  $f_{\tau}$  is dominated by the transit delay in the device while  $f_{max}$  is approximately  $\propto C_{cb}^{-1}$ , and digital logic speed is severely impacted by the delay term,  $C_{cb} \Delta V_{logic}/I_c$ . The numbers in Table 2.2 serve to illustrate that reduction in  $C_{cb}$  considerably enhances the high frequency performance of power amplifiers (which depends on  $f_{max}$ ), and digital circuits.

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# 3

## Ion Implantation in InP

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ION implantation is a statistical process, where high energy ions penetrate a target surface and are slowed down in traversing matter. The ions of a material are implanted into another solid changing the physical properties of the solid. The ions impinge on the substrate with kinetic energies 45 orders of magnitude greater than the binding energy of the solid substrate.

Ion implantation is the method of choice in state of the art Si manufacturing to bring the dopants into the substrate material, mainly due to its ability to accurately control the number of implanted dopants and to place them at the desired depth. Ion implantation works by ionizing the required atoms, accelerating them in an electric field, and directing this beam towards the substrate. When entering the substrate material the energy of the dopants decreases, while they interact with the target material. After some time the atoms come to rest at some depth depending on their initial energy. This depth has some distribution as the collisions with the target atoms are random. An important point for the device design is to know which initial energy is necessary to place the dopants at the required depth and what will be their spread. The LSS (Lindhard, Scharff, Schitt) theory [1] describes the distribution of the ions in matter based on statistical models of atom-atom collisions .

### CHAPTER 3. ION IMPLANTATION IN INP

The two key parameters defining the final implant profile are fluence, also called as dose, given in ions/cm<sup>2</sup> and energy E (in keV ). The dose or the areal flux of the ions, is related to the beam current I by the following formula:

$$\phi = \frac{I \cdot t_I}{q_i \cdot A_i} \quad (3.0.0)$$

where  $t_I$  denotes implantation time,  $A_i$  the beam area and  $q_i$  is the charge per ion. Typical beam currents and implantation doses range from 1  $\mu$ A- 30mA and  $\times 10^{11}$  -  $\times 10^{16}$  ions/cm<sup>2</sup>. The lowest energies used start at sub keV for ultra shallow junctions to the MeV range for deep wells. When the ions enter the substrate they continuously lose energy and momentum. There are two main effects that causes an energy loss,

- elastic collisions with the nuclei of the target material
- inelastic collisions with the electrons

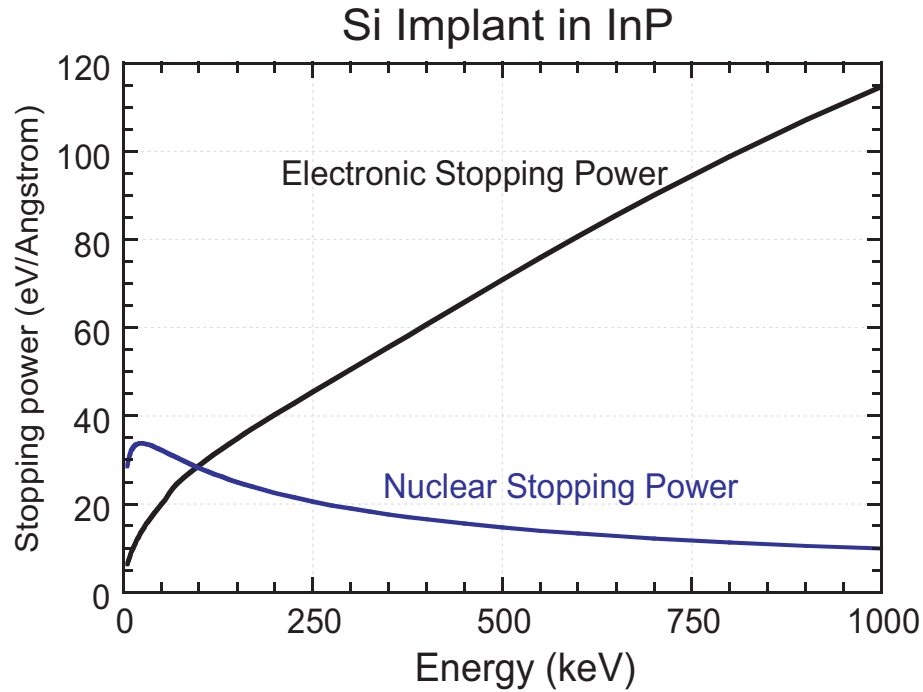


Figure 3.1: Nuclear and Electronic Stopping Power of Si implant in InP

The total stopping power S defined as the energy loss per unit path length of the ion can be defined as:

$$S = \frac{dE_{ii}^{nuclear}}{dx} + \frac{dE_{ii}^{electronic}}{dx} \quad (3.0.0)$$

### CHAPTER 3. ION IMPLANTATION IN INP

where  $E_{ii}$  is the initial ion energy on impact.

$$\frac{dE_{ii}^{electronic}}{dx} \propto \sqrt{E_{ii}} \quad (3.0.0)$$

Both these components are functions of the mass of the target and incident ion and nuclear charge of the target. Fig. 3.1 is a plot of electronic and nuclear stopping powers calculated from *TRIM*, for Si implants in InP. The nuclear stopping is dominant at low energies while at high energies, the stopping is almost entirely due to the electrons of the target material, InP.

Due to the random nature of the collisions the total distance traveled (range) and its projection on the direction parallel to the ion beam (projected range) are random variables.  $R_p$  denotes the *projected range*, the depth where most ions stop. The *longitudinal projected straggle*  $\Delta R_p$ , describes the statistical fluctuation of  $R_p$ . Based on the LSS theory, the implant profile (projected ranges  $R_p$  of a huge number of ions) is described by a Gaussian function since the sum of infinite random variables can be approximated by a Gaussian distribution as,

$$N(x) = \frac{n_{ii}}{\Delta R_p \sqrt{2\pi}} \cdot \exp - \left( \frac{(x - R_p)^2}{2\Delta R_p^2} \right) \quad (3.0.0)$$

where  $n_{ii}$  is the implant fluence in ions/cm<sup>2</sup>,  $R_p$  is the implant range and  $\Delta R_p$  is the longitudinal implant straggle. The profile is defined by the implanted fluence  $n_{ii}$  in ions/cm<sup>2</sup>, the projected range  $R_p$  in  $\mu\text{m}$  and the projected straggle  $\Delta R_p$  in  $\mu\text{m}$ .

Thus, the range and straggle are functions of the implanted species, initial energy, and target material. Since each ion's final distribution is a random variable, statistical Monte Carlo simulations can predict the projected range and straggle of a huge number of ions. The software SRIM [2] is a group of programs which calculate the stopping and range of ions into matter using a full quantum mechanical treatment of ion-atom collisions. One component *TRIM* (the Transport of Ions in Matter) is a Monte Carlo Transport Calculation of ion interactions with multi-layer complex targets. It can calculate both the final 3-D distribution of the ions and also all kinetic phenomena associated with the ion's energy loss: target damage, sputtering, ionization, and phonon production while all target atom cascades in the target are followed in detail.

*TRIM* assumes that there is cylindrical symmetry in the final ion distributions. Thus the mean lateral range of the ions is zero while the ion straggling has its normal definition as the second moment of the lateral distribution. The lateral projected straggle thus directly describes the final distribution under an implant mask edge.

### 3.1 Implantation in InP

Seminal work on implantation in InP was carried out in the late 70s and early 80s by Donnelly and Hurwitz [3]. They demonstrated that Ion implantation is a feasible method to achieve either doping or isolation in InP. The magnitude of resistivity of implanted InP depends on the doping species. It is seen that activating acceptor implants are harder than donor implants. Another problem with acceptor species is their high rate of diffusion in InP. The implanted species Si, Ge, Se, S and Sn are all effective donors in InP and can produce heavily doped N-type layers. Many electronic applications require Semi Insulating(SI) InP with high resistivity of  $\sim \times 10^{10} \Omega \cdot \mu\text{m}$ . Resistivities in the SI range are usually obtained by doping with the deep acceptor Iron(Fe) or as is done in this thesis, by implanting Fe. All these implants require high temperature annealing at over 700°C.

As the implanted ions interact with the target nuclei, they can initiate a cascade of displacements. In general, the heavier ions create more damage. At high substrate temperature, defects are more mobile and more annealing can take place during the implant. It is seen in InP that there is significantly less damage if the implants are carried out at an elevated temperature over 170 °C [4]. Furthermore, the activation of the dopants and the mobility significantly improve when the implants are carried out at elevated temperatures [5]. This is attributed to a dynamic annealing process during implant. Since the implants are carried out at an elevated temperature, an implant mask should be used, that can withstand these temperatures.

During implantation, channeling can occur when the ion velocity is parallel to a major crystal orientation. The ions can travel considerable distances with little energy loss. Ion channeling can therefore produce a significant tail to the implant distributions. Most IC implantation is therefore done off-axis. A typical tilt angle is 7°. Channeling is more a problem in single crystal materials than in III-V compounds. The power of an incident ion is

$$P_{ii} \propto V_{ii} \times J_{ii} \quad (3.1.0)$$

where  $J_{ii}$  is the ion current flux in  $\text{mA}/\mu\text{m}^2$  and  $V_{ii}$  is the voltage =  $E_{ii}/\text{ioncharge}$ . If the incident power is transferred to the target crystal, it is dissipated as heat. In typical VLSI processes, where the implants in Si are carried out at room temperature or for damage implants in III-V's, this temperature rise can destroy the photoresist typically used as an implant mask. Therefore current is usually maintained small in such cases, usually  $\leq 50 \mu\text{A}$  which means that for a given implant fluence, the implantation times can be very long and hence there is a higher probability of impurities. All implants done here are at a substrate temperature of 200 °C. Therefore, the implant current is less important for such implants. However, to not cause additional lattice heating, all implants are carried out at  $\leq 50 \mu\text{A}$ .

### 3.1.1 High temperature annealing of implants in InP

One component of energy transfer when a high energy ion enters a wafer is collision with the lattice nuclei. Many of these atoms are ejected from the lattice during the process. Some displaced substrate atoms have sufficient energy to collide with other substrate atoms to produce additional displaced atoms. As a result, implantation process can produce considerable substrate damage that must be repaired during subsequent processing. Furthermore, if the implanted species is intended to act as a dopant, it must occupy lattice sites, here group III In sites. Both damage repair and implant activation are normally done by heating (annealing) the wafer after implant.

There exists a threshold dose above which the damage is complete. This critical dose depends on the implant energy, implant and target species and substrate temperature. As an ion passes through the crystal, point defects consisting of interstitials and vacancies are created by direct interactions with the target atoms. These are primary defects. A vacancy defect is also known as Schottky defect and an interstitial defect is called a Frenkel defect. Secondary defects occur when an implanted wafer is annealed. Secondary defects increases greatly at very high fluences near the critical dose [6].

Annealing processes are needed that minimize the secondary defects while activating the dopants. The temperature required for optimum activation of dopants in InP is well above the temperature (550°C) at which incongruent evaporation of group V species, Phosphorus P, occurs. Therefore some method of minimizing this loss is required. A common method of annealing is the proximity method where another InP wafer is placed face to face with the implanted InP substrate. As the wafers are heated up, each wafer begins to lose a small amount of P, but an overpressure is created that prevents further dissociation. This is not a satisfying solution since it relies on loss of P. Moreover it presents a safety issue when a conventional RTA chamber is used since P is highly flammable and poisonous. The best method is to anneal the InP wafer in a phosphorus atmosphere in the MBE chamber, but this choice is not available at this time at UCSB. Annealing in a enclosed graphite cavity with a P partial pressure is shown to be very effective even upto 900 °C [7]. A commonly used solution is the use of dielectric encapsulant such as  $SiO_2$ ,  $Si_xN_y$  or Phosphosilicate glass (PSG) [3] where the wafer can be annealed in a conventional RTA.

$SiO_2$  is a poor encapsulant as it is hard to remove after anneal and also has vastly different thermal expansion coefficient from InP.  $Si_xN_y$  is used as an encapsulant here. In order to minimize scratching on the  $Si_xN_y$  on the top surface during backside deposition, the PECVD platen is first coated with 100-200 nm of  $Si_xN_y$ . All anneals are carried out in the Rapid Thermal Annealer (RTA) as this was the only tool available at this time. Furthermore, if the anneal is carried out at high tempera-

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ture and for a short time, thermal equilibrium is not reached. Thus dopants greater than its solid solubility in InP can be activated. Stress and strain can cause cracking or a loss of adhesion, so it is important to use pinhole free  $Si_xN_y$ . Both sides of the wafer are coated with 40 nm of  $Si_xN_y$  deposited by PECVD. The quality and thickness of  $Si_xN_y$  are very critical to providing surface integrity during anneal. For annealing, the deposited  $Si_xN_y$  has to be very uniform and very smooth without

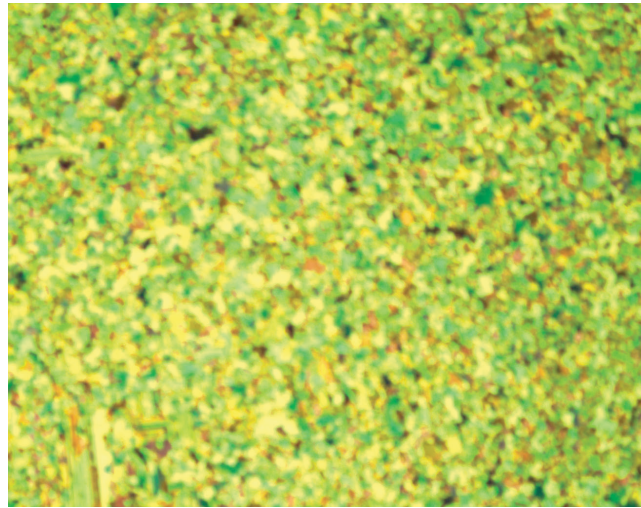


Figure 3.2: Poor quality  $Si_xN_y$  on InP

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pinholes or defects. An example of poor quality  $Si_xN_y$  on InP is seen in Fig. 3.2.  $Si_xN_y$  has poor adhesion on InP and often results in a rough layer. Deposition of good quality  $Si_xN_y$  on InP is not trivial. PECVD deposited  $Si_xN_y$  adheres poorly to exposed InP surfaces. Prior to deposition, InP surfaces are treated with Ozone for 5-10 minutes. Studies by Driad *et al.* [8] showed that an ultra-violet ozone treatment (UV-ozone) is an effective way of cleaning a surface of organic and non-organic materials. The uv-Ozone also produces a uniform stoichiometric oxide film which passivates the defective surface layers associated with the native oxides of InP and those that occur due to processing. After a ten minute uv-Ozone treatment, the stoichiometric composition at the surface is restored. The oxide thus formed on InP has to be removed prior to  $Si_xN_y$  deposition. This is done by wet etching in Buffered Hydro-Fluoric(BHF) acid for 5-10 minutes, so that a clean surface of InP ensues. However, Fluorine radicals from BHF are attracted to InP surface [9]. Therefore the wafer is cleaned in running De-Ionized water (DI) for 5-10 minutes. It is seen that each of these steps is critical in ensuring a clean InP surface and hence good quality  $Si_xN_y$  as cap.

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The thickness of the  $Si_xN_y$  cap is important. A thin layer of  $Si_xN_y$   $\leq 30$  nm may not provide adequate surface coverage and is not reliable. A thicker  $Si_xN_y$  cap  $\geq 60$  nm results in numerous defects after anneal. The reasons are not well understood at this time. One postulate is that the  $Si_xN_y$  layer is strained and a thick cap may be relieving this strain at high temperatures leading to pinholes. The pinholes results in P desorbing from the surface and leaves behind deep pits  $\sim 15 - 20 \mu\text{m}$  as seen in Fig. 3.4. Fig. 3.3 is such a micrograph of such a defect on the InP surface after a high temperature anneal(800 °C). Analysis of this anneal defect using Energy Dispersive X-Rays (EDX) indicates a very large concentration of Oxygen atoms. The atomic ratio of P to In is slightly lower.

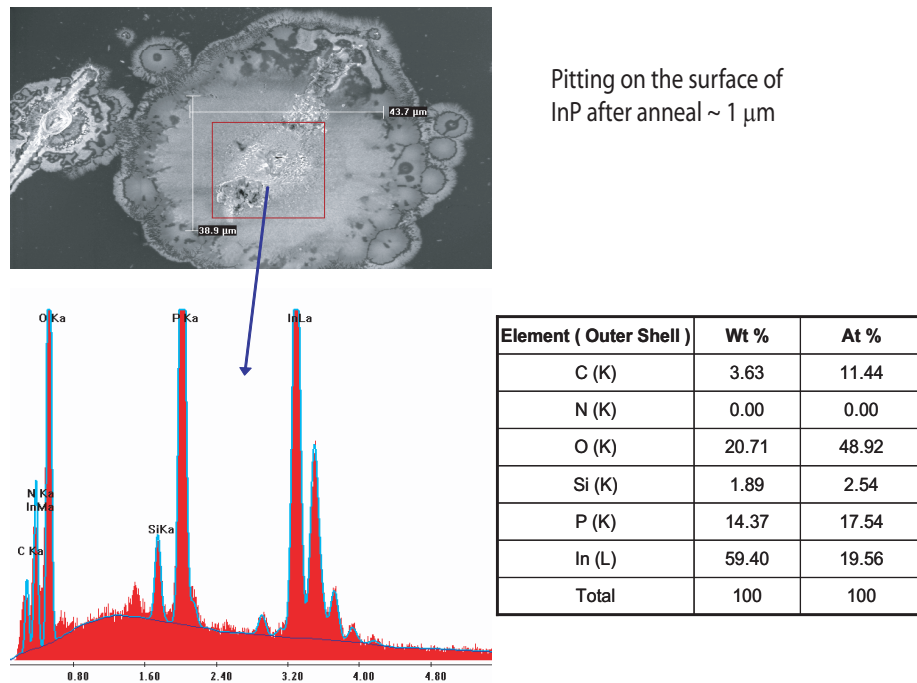


Figure 3.3: Rectangular pit formation on InP after high temperature anneal and EDX compositional analysis

The complete removal of  $Si_xN_y$  after anneal is critical. High temperature anneals cause the dielectric encapsulant to become stoichiometric and dense. These are typically harder to remove after such high temperature treatments. This was the main reason for rejecting  $SiO_2$  as an encapsulant. It is seen that a long treatment in BHF ( $\sim 10$  minutes) is sufficient to remove this  $Si_xN_y$  cap. Fig. 3.5 is an EDX spectra of the InP surface after the  $Si_xN_y$  is removed in BHF and there is no



### CHAPTER 3. ION IMPLANTATION IN INP

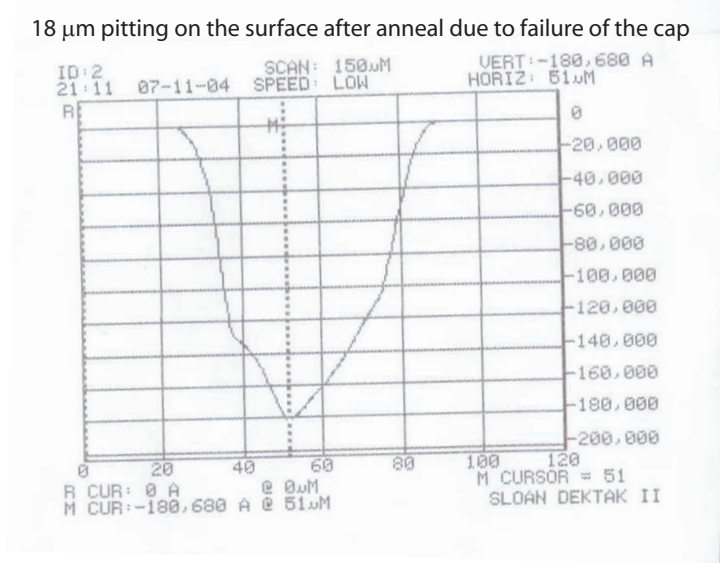


Figure 3.4: Pit formation on InP after high temperature anneal

evidence of  $Si_xN_y$  present.

One important consideration is the residual stress of the encapsulant [10]. Compressive stress may cause undesired buckling and cracking may occur if the tensile stress is too high. The bulk stress consists of an intrinsic and thermal part. The total bulk stress ( $\sigma_{bulk,total}$ ) in dielectric PECVD layers is given by

$$\sigma_{total} = \sigma_{int} + \sigma_{th} \quad (3.1.0)$$

where  $\sigma_{int}$  and  $\sigma_{th}$  are the bulk intrinsic and thermal stress respectively. The thermal stress only results from the temperature change between deposition and anneal and the difference in thermal expansion coefficients of the film and the substrate. The thermal stress is given by

$$\sigma_{th} \propto (\alpha_f - \alpha_s)\Delta T \quad (3.1.0)$$

where  $\alpha_f$   $\alpha_s$  are the thermal expansion coefficients of the film and substrate respectively. An encapsulant should therefore ideally have a thermal expansion coefficient similar to InP to prevent thermal stresses. Between 300 and 673K, the thermal expansion coefficient of InP is  $4.6 \times 10^{-6}/^\circ K$  while that of  $Si_xN_y$  is  $\sim 3.2 \times 10^{-6}/^\circ K$ .  $SiO_2$  has a thermal expansion coefficient of  $0.5 \times 10^{-6}/^\circ K$  which makes it less suitable as an encapsulant for InP.

In the long term however, annealing of InP in a phosphorus overpressure in a fully sealed chamber seems to be the best solution.

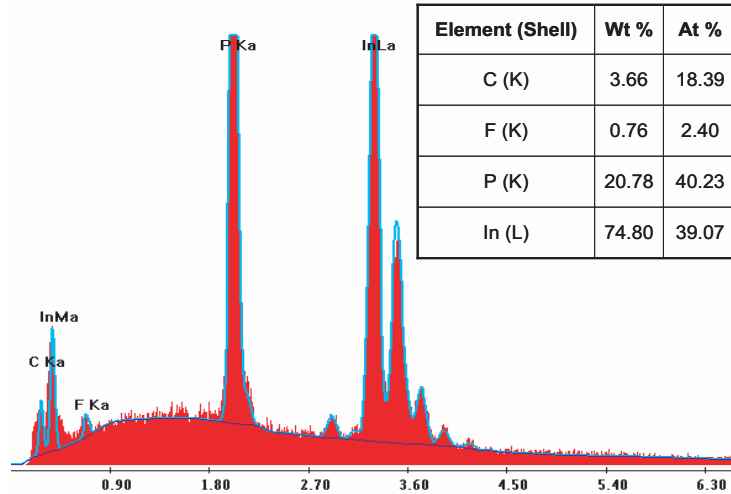


Figure 3.5: Analysis of the InP surface after anneal and cap removal

### 3.1.2 Si implantation in InP

N-type dopants implanted in InP have been shown to have an activation  $\geq 80\%$  even when the doping levels is  $\sim 1 \times 10^{19} \text{ cm}^{-3}$  [11]. Si, Se and S are commonly used dopants. Among these Si is the lightest element and causes minimum damage. It has little diffusion which is not the case with S [12] and therefore the lateral spread can be better controlled with Si implants. Here, Si implantation is done wherever N-doping is required. The commercially available ion implant systems available at this time could do a maximum of 350 keV at a substrate temperature of 200 °C. Fig. 3.6 shows the distribution of Si in InP, as estimated by TRIM, when implanted at 350 keV. TRIM does not account for differences due to implant temperatures. Also this plot is a plot of ion range in  $\text{atoms}\cdot\text{cm}^{-3} / \text{atoms}\cdot\text{cm}^{-2}$ . When multiplied by the dose in  $\text{ions}/\text{cm}^2$  or ( $\text{atoms}\cdot\text{cm}^{-2}$ ) it gives the distribution of Si in  $\text{cm}^{-3}$ .

The Ion recoil distributions are seen in Fig. 3.7a. These are all the In and P atoms knocked out of their lattice sites, creating vacancies. The distribution of In and P recoils is shallower than the range of the Si (Fig. 3.6). Near the end of the Si tracks, the ions do not have enough energy to create massive cascades. At the peak of the damage plot in Fig. 3.7a, the vacancy rate of In is  $\sim 0.4$  vacancy/ion-Å. For a dose of  $5 \times 10^{14} \text{ ions}/\text{cm}^2$ , there are  $2 \times 10^{22} \text{ vacancies}\cdot\text{cm}^{-3}$ . The atomic concentration of In is  $5.6 \times 10^{22} \text{ atoms}\cdot\text{cm}^{-3}$ . The damage is therefore  $\sim 35\%$ . This is a pessimistic estimate. Due to elevated substrate temperature,  $\sim 90\%$  of the damage is repaired

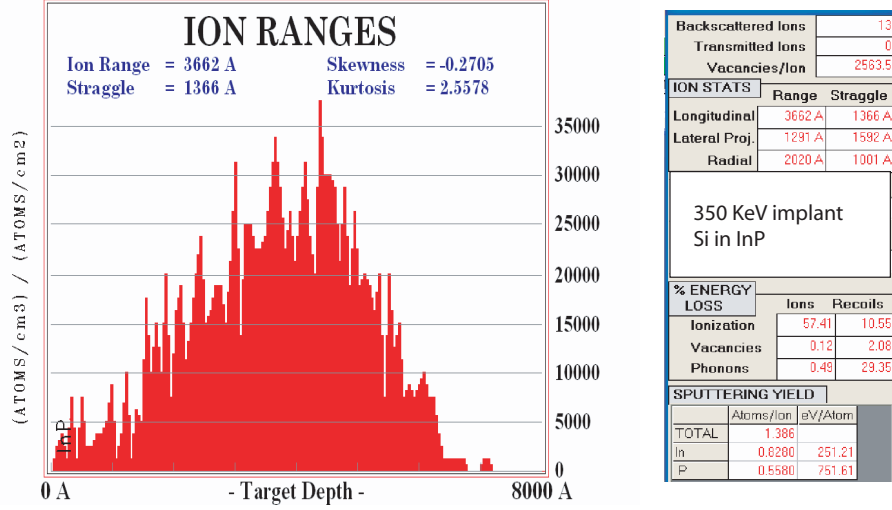


Figure 3.6: Ion Ranges for Si in InP at 350 keV

during implant. Therefore the InP crystal damage is  $\sim 3.5\%$  and the implant layers are not quite amorphous.

Fig. 3.7b indicates the energy lost to recoiling target atoms that give rise to vacancies. From the plot,  $\sim 200$  keV of the ions energy is transferred to the recoiling In atoms. This can be estimated by assuming a constant energy transfer of  $\sim 25$  eV/ion- $\text{\AA}$  over a depth of  $8000\text{\AA}$ . So the total energy transfer to In recoils is  $\sim 200$  keV. Also, it is seen that the energy transfer to phonons is almost exclusively from the recoiling target atoms. An atom is knocked off its lattice site, giving rise to a vacancy when the energy is greater than the lattice binding energy. The total displaced atoms from their lattice sites is usually quantified as damage due to implant. These atoms leave behind vacancies and may form interstitials.

The Nuclear stopping power and electronic stopping power is also calculated from *TRIM*. At 350 keV the stopping power in eV/ $\text{\AA}$  is,

$$S_n = \frac{dE_{ii}^{nuclear}}{dx} = 17.69 \text{ eV}/\text{\AA} \quad (3.1.1)$$

$$S_e = \frac{dE_{ii}^{electronic}}{dx} = 55.65 \text{ eV}/\text{\AA} \quad (3.1.2)$$

From the projected range and longitudinal straggle obtained from *TRIM*, and assuming a Gaussian distribution, the implanted profile of Si is plotted in Fig. 3.8. Also shown are the doping levels with 60 % and 100 % activation of the Si dopants.

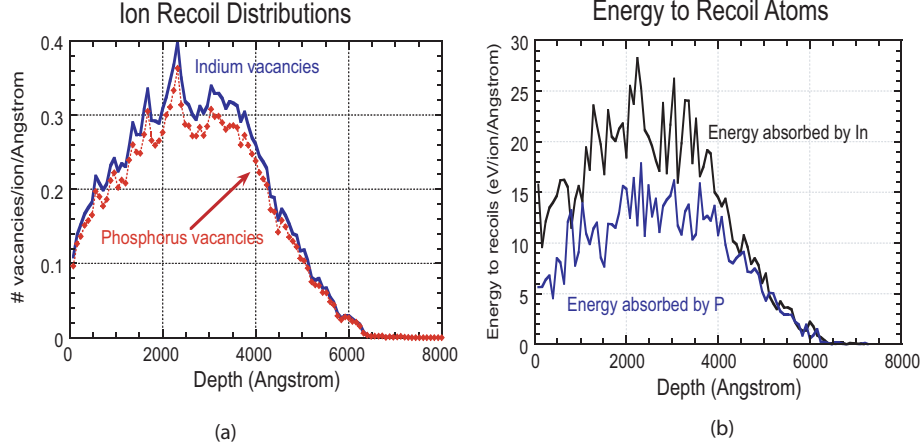


Figure 3.7: Ion Ranges for Si in InP at 350 keV

With 60 % activation, the average doping is  $\sim 5 \times 10^{18} \text{ cm}^{-3}$  over 6000 Å, and using a mobility of  $\sim 1000 \text{ cm}^2/\text{V} \cdot \text{s}$ , the expected sheet resistance is  $\sim 21 \Omega/\square$ .

The lateral straggle is given as 1592Å. As discussed, cylindrical symmetry is used. Since the offset angle is small, the lateral range is assumed to be zero. The lateral distribution can be expressed as,

$$N(z) = N(x) \cdot \exp - \left( \frac{z^2}{2\Delta R_z^2} \right) \quad (3.1.2)$$

where  $R_z$  is the lateral straggle distance. Lateral straggle is the distance in  $z$  when the concentration reduces to  $e^{-1/2}$  of its peak value. The peak activated value (at the implant mask edge) is a function of  $x$  as seen in Fig. 3.8 and is  $\sim 1 \times 10^{19} \text{ cm}^{-3}$ . The lateral distribution is plotted in Fig. 3.9. The point at which the Si doping falls below the Fe doping present in SI InP, the region become Semi-Insulating again. In this dissertation, this is important and is defined as the lateral straggle due to Si implant. This straggle is  $\sim 0.5 \mu\text{m}$ . Table 3.1 lists the projected range and longitudinal and lateral straggle for Si at various energies in InP from Monte Carlo simulations.

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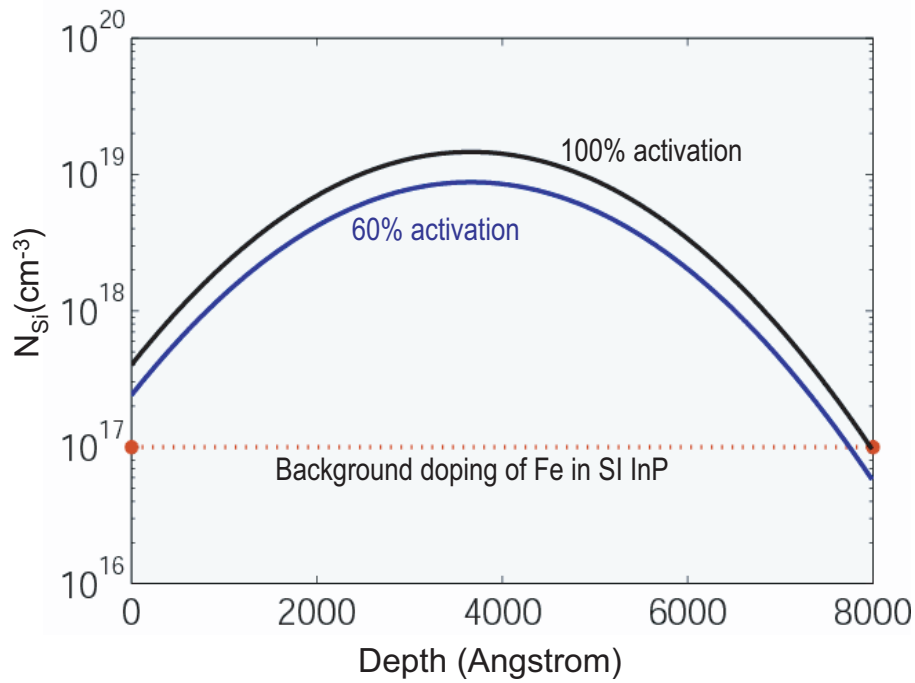


Figure 3.8: Si doping profile in SI InP at 350 keV

Table 3.1: Silicon Implant in InP as calculated by TRIM

Implant energy	Projected Range	Longitudinal Straggle	Lateral Straggle
5	92	52	54
10	159	91	85
40	496	256	286
90	1020	490	540
140	1556	696	800
200	2204	902	1060
350	3662	1366	1592

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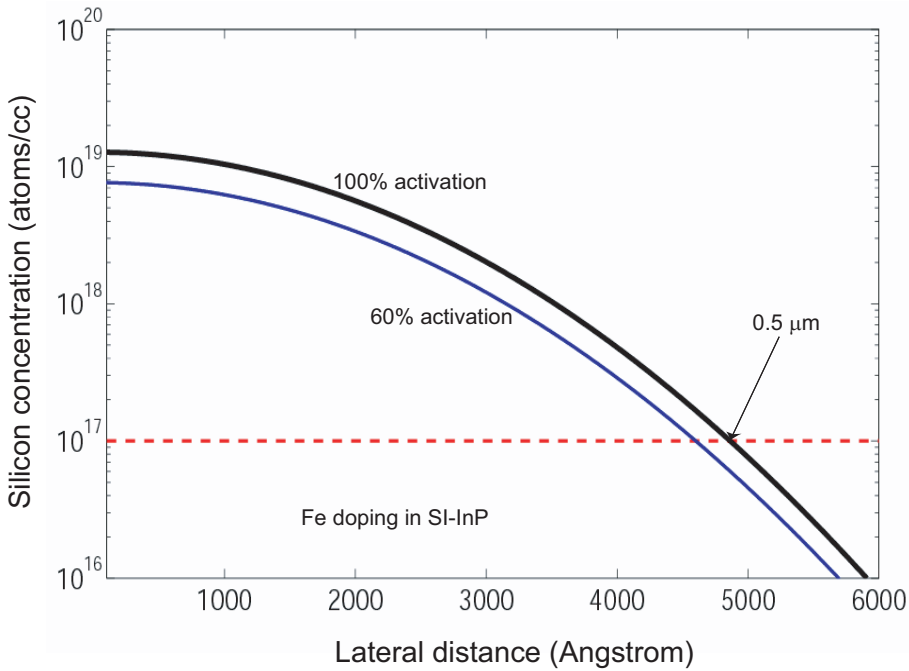


Figure 3.9: Lateral straggle of the 350 keV Si implant

### CHAPTER 3. ION IMPLANTATION IN INP

Table 3.2: Implant conditions for basic, implanted subcollector HBT process

Implant species	Implant energy	Implant fluence	Offset angle	Implant temperature
Si	140 keV	$3 \times 10^{14}$ ions/cm <sup>2</sup>	7 °	200 ° C
	40 keV	$8 \times 10^{13}$ ions/cm <sup>2</sup>		
	10 keV	$3 \times 10^{13}$ ions/cm <sup>2</sup>		
$I_{ions} \leq 10 \mu A$				

Si implants are performed at the implant conditions stated in Table 3.2. After implant, the wafers were capped with  $Si_xN_y$  and annealed at various temperatures and times. Table 4.7 gives the mobilities and sheet charge density. It has been

Table 3.3: Hall measurements of resistivity and mobility for Si implants

Anneal temperature (°C)	Anneal time (s)	Hall mobility (cm <sup>2</sup> /V · s)	Carrier density (cm <sup>-2</sup> )	Sheet resistance (Ω/□)
750	30	999	$1.22 \times 10^{14}$	51.3
800	15	868	$1.70 \times 10^{14}$	42.3
800	30	909	$1.66 \times 10^{14}$	41.5
825	15	874	$1.87 \times 10^{14}$	38.3
850	15	879	$2.11 \times 10^{14}$	33.7

reported [11] that short anneals at temperatures over 775 °C is required for complete activation of donor implants. However, above 800 °C the  $Si_xN_y$  cap is not stable and there are numerous anneal defects throughout the wafer. The activation of Si for 800°C anneal is ~60%. Phosphorus(P) co-implantation is known to greatly increase the activation of Si. Si acts as donor only if it substitutes *In* atoms in the lattice. Co-implanting P increases this possibility. The profile of Si implant is obtained from SIMS and is shown in Fig. 3.10. It deviates little from the *TRIM* simulations.

The sheet resistance obtained above is high for the applications discussed in later chapters. Therefore the implant energy is increased, to enhance the implant depth. The details are in Chapter 4 . AFM scans in Fig. 3.11 of the implanted and annealed wafers indicate that the surface quality is restored to its original state after annealing at 800 °C.

CHAPTER 3. ION IMPLANTATION IN INP

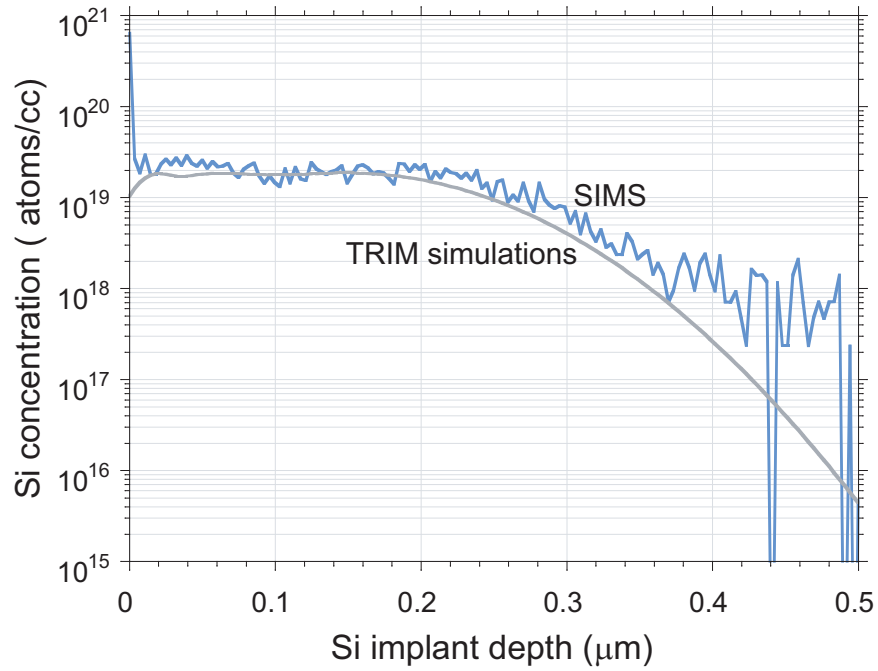


Figure 3.10: Si doping profile in InP at 140 keV from SIMS and simulated by *TRIM*

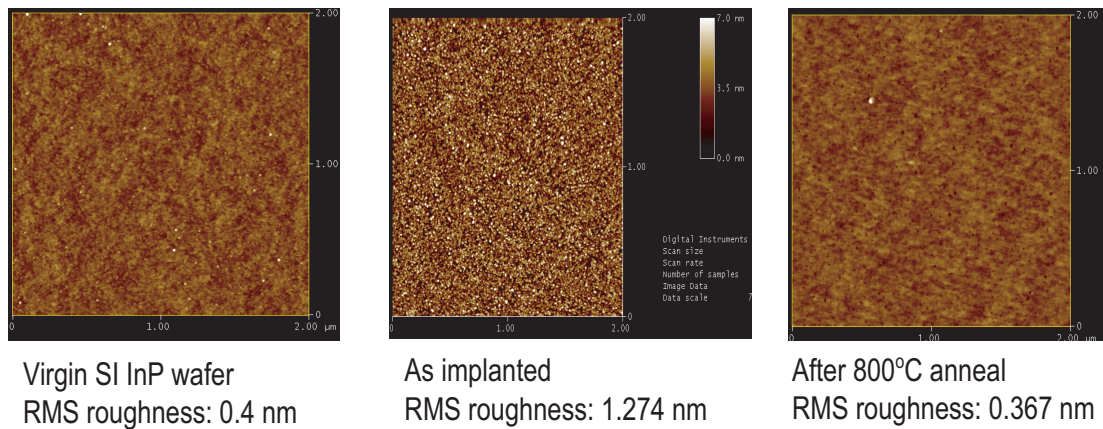


Figure 3.11: AFM scans of the implanted surface



### 3.1.3 Fe implantation in InP

Iron (Fe) is a mid gap acceptor in InP. Fe is a transition metal with tightly bound orbitals and hence a small Bohr Radius. Fe atoms occupy the In sites substitutionally and act as deep acceptor centers for free electrons. Indeed, SI InP substrates are formed by doping the crystal, during the Liquid Encapsulated Czochralski (LEC) growth phase, with Fe. Fe typically exists in InP with an oxidation state of +3 and it can accept one electron to reduce its oxidation state of +2.

Isolation can also be done by implantation. However in InP, implant isolation has not been as effective in creating high resistivity regions as in GaAs. The ion species H, He, B, O<sub>2</sub> are utilized to create damage implants in n-type InP and, the reported value of 1 MΩ/□ is insufficient for device isolation. Fe Implantation can be used to create deep levels; this chemical compensation ensures the crystallinity of the semiconductor after high temperature anneal. Isolation by chemical compensation is more apt if there is a requirement of subsequent regrowth of crystalline layers. Full compensation is achieved when the concentration of the deep acceptor levels  $N_A$ , exceeds the background doping  $N_D$ , the Fermi level  $E_F$ , is approximately pinned at  $E_A$ , the deep acceptor level due to Iron which is  $\sim 0.6\text{eV}$  below the conduction band [13]. This result is derived as follows. Consider a semiconductor with a shallow donor of active density  $N_D$  and a deep acceptor at  $E_f$  and active density  $N_A$ .  $n$  is the number of conduction band electrons and  $p$  is the number of valence band holes. Assuming Boltzmann statistics,

$$n = N_C e^{-(E_C - E_F)/kT} \quad (3.1.3)$$

$$p = n_i^2/n \quad (3.1.4)$$

where  $n_i$  is the intrinsic concentration in the semiconductor. The Fe acceptor has a negative charge if occupied and neutral if empty. The occupation of acceptors is given by,

$$n(E_A) = \frac{N_A g_A}{g_A + e^{-(E_F - E_A)/kT}} \quad (3.1.5)$$

$$p(E_A) = N_A - n(E_A) \quad (3.1.6)$$

where  $g_A$  is the degeneracy of the acceptor states.  $g_A$  is usually 4. Charge neutrality dictates that,

$$n + n(E_A) = p + N_D^+ \text{ where, } p = n_i^2/n \quad (3.1.6)$$

Complete ionization of the donor states is assumed.  $n(E_A)$  can be simplified as,

$$n(E_A) = \frac{N_A}{1 + \theta/n} \text{ where,} \quad (3.1.7)$$

$$\theta = \frac{N_C}{g_A} \exp \frac{E_A - E_C}{kT} \quad (3.1.8)$$

### CHAPTER 3. ION IMPLANTATION IN INP

The charge neutrality equation can therefore be written in simplified form as,

$$n + \frac{N_A}{1 + \theta/n} = N_D^+ + \frac{n_i^2}{n} \text{ or,} \quad (3.1.9)$$

$$\frac{n^2}{N_D^2} + \left( \frac{N_A/N_D}{1 + \theta/n} - 1 \right) \frac{n}{N_D} - \frac{n_i^2}{N_D^2} = 0 \quad (3.1.10)$$

If  $N_A/N_D \geq 1$  and the semiconductor become semi-insulating, or intrinsic, then the number of conduction electrons is determined by the thermal generation, or  $n = n_i$ . In that case,

$$\frac{N_A/N_D}{1 + \theta/n} = 1 \text{ or,} \quad (3.1.11)$$

$$n = \frac{\theta}{x - 1} \text{ where, } x = \frac{N_A}{N_D} \quad (3.1.12)$$

or expanding  $\theta$  and  $n$  from Eqn.3.1.4 and Eqn.?? one obtains,

$$E_F = E_A - kT \ln[g_A(x - 1)] \quad (3.1.12)$$

As mentioned, this expression is only valid when  $x = \frac{N_A}{N_D} \geq 1$ . A plot of the Fermi level with  $x$  is given in Fig. 3.12. As evident from the figure, the Fermi level is pinned at the acceptor level even when  $x = 20$ . This is the reason why mid gap acceptors are used to compensate residual donors, when a SI semiconductor are needed. When the Fermi level is above the acceptor level, the acceptor states are filled. Since all the acceptor states are below the donor energy levels, the electrons preferentially occupy the acceptor sites. This means that the donor states are ionized and, if the donor density is less than the acceptor density, the conduction band is  $\approx$  empty. In this case the Fermi level is at  $\approx E_A$ . Suppose more donors are added. First, all the acceptor states are completely filled. The remaining donors then ionize so that their electrons move into the conduction band. In this case, the Fermi level approaches the donor level as seen from Fig. 3.12.

The solid solubility of Fe in InP is  $1 \times 10^{17} \text{ cm}^{-3}$ . However, this can be overcome during implantation, since it is essentially a non equilibrium process. Donnelly and Hurwitz first reported the effect of iron bombardment in InP. Since then, numerous studies have been conducted using iron implantation in InP [13, 14, 15]. Earlier work indicates [11] that prolonged anneal at temperatures over  $650 \text{ }^\circ\text{C}$  is required for damage removal and complete activation of acceptor implants. Indeed, resistivities as high as  $2 \times 10^7 \text{ } \Omega\text{-cm}$  have been obtained, for Fe implanted in  $N^{++}$  doped InP ( $1 \times 10^{14} \text{ cm}^{-3}$ ), after  $650 \text{ }^\circ\text{C}$  anneal. However, room temperature implantation of Fe produces an amorphous implant region which results in significant defects even after anneal [16].

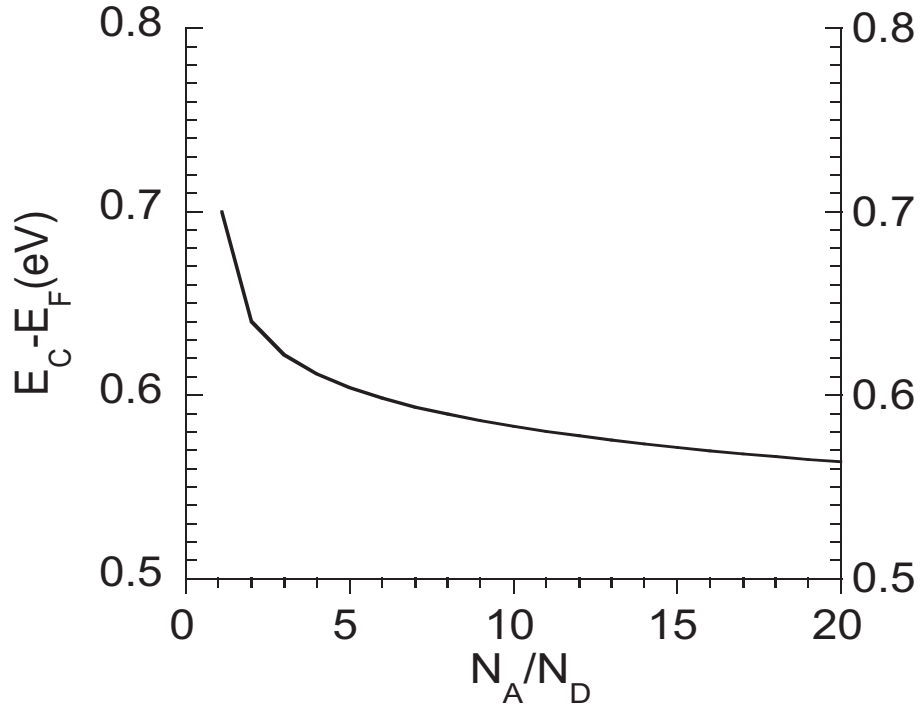


Figure 3.12: Fermi levels for various degrees of deep level compensation

To study the effect of Fe in  $N^{++}$  InP, Fe is implanted at 190 keV with a fluence of  $4 \times 10^{15}$  ions/cm<sup>2</sup> into epitaxially grown 130 nm  $N^-$ InP and 80 nm  $N^{++}$  InP. This was carried out at 200 °C to minimize crystal damage and increase Fe activation, which occurs due to enhanced dynamic annealing at this temperature [17]. As before, the post implant is carried out in a conventional a conventional RTA. Fig. 3.13 shows the SIMS profiles of Fe in the as-implanted and annealed wafers and also the theoretical LSS profile from *TRIM*. There is significant deviation from the theoretical profile, in that

- the Fe penetration is deeper than predicted by the TRIM software
- a reduction in the peak concentration
- a pileup of Fe at the surface

. These results concur with that reported earlier [18, 19, 15]. Several arguments exists to explain this phenomena. Fe tends to accumulate near the maximum damage

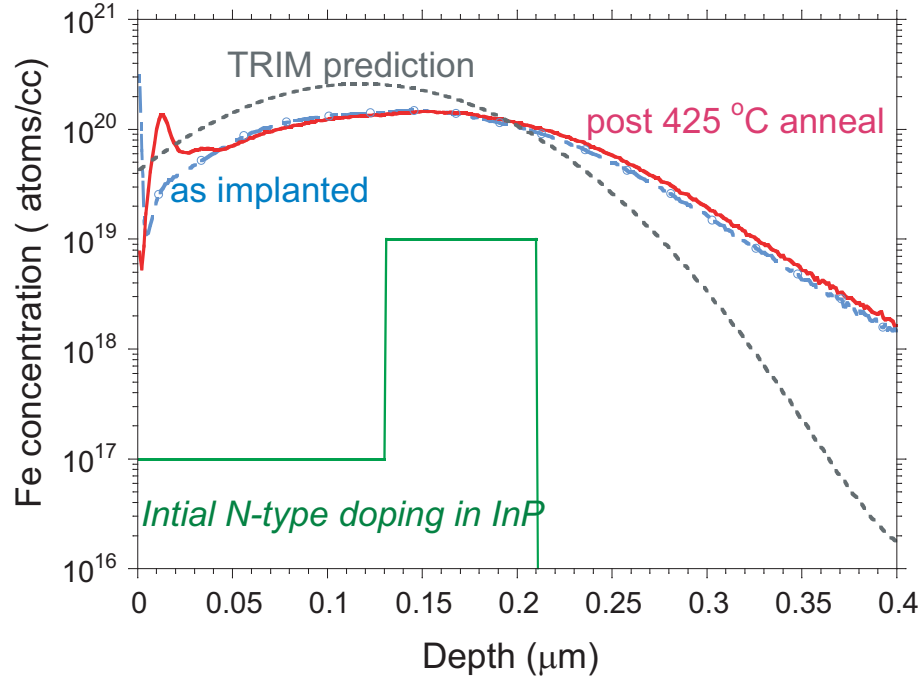


Figure 3.13: Fe profile in 130nm  $N^-$ /80nm  $N^{++}$  InP at 190 keV/ $4 \times 10^{15}$  ions/cm<sup>2</sup> from SIMS and as simulated by *TRIM*

region with peaks at around  $R_p + \Delta R_p$  of the Fe implant. This discrepancy is explained by the enhanced diffusivity of Fe in the amorphous phase [20]. The diffusion coefficient is  $\sim 1 \times 10^{13}$  cm<sup>2</sup>s<sup>-1</sup>. Furthermore such accumulation at certain points, occur when the Fe implant fluence exceeds a certain amorphization threshold. This threshold is  $1 \times 10^{13}$  ions/cm<sup>2</sup> when implanted at room temperature [20] and between  $2$  and  $5 \times 10^{14}$  ions/cm<sup>2</sup> when implanted at 200 °C. A depletion of Fe atoms is also observed and this larger when the damage is higher. The Fe atoms are gettered and/or trapped at the secondary defects formed during annealing. Reconstruction of an amorphous layer is complicated and strongly influences the redistribution properties of Fe. . The RBS plots in Fig. 3.14 indicate the presence of an amorphous region upto a depth of 104 nm for the implant performed at 190 keV with a fluence of  $4 \times 10^{15}$  ions/cm<sup>2</sup>. The anneal at 425°C is not sufficient to repair the crystal damage. As evident from SIMS, the implanted Fe concentration is well over the Si N-type doping in the structure. Fig. 3.15 shows a plot of measured resistivity of the sample with anneal temperature; the as-implanted sheet resistance is 3 MΩ/□. At temperatures of 500°C and above, the resistance of the Fe implanted layers drops.

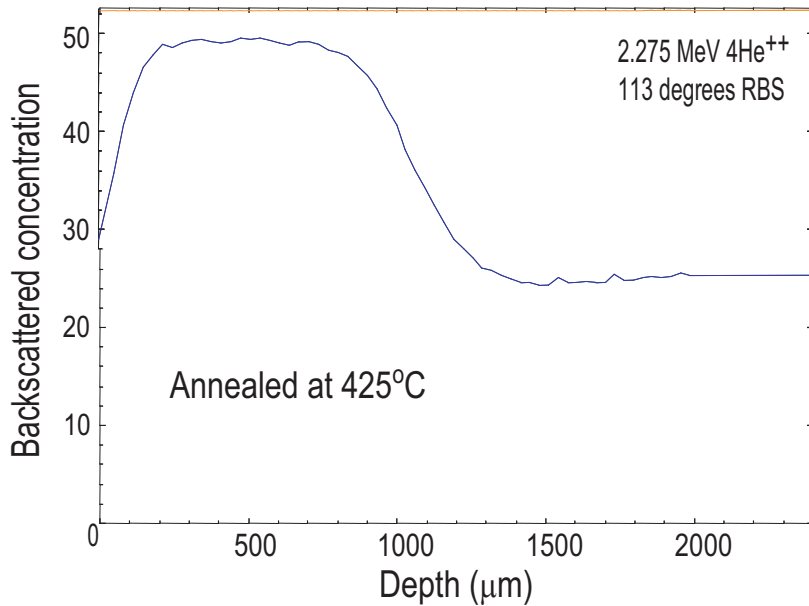


Figure 3.14: RBS plots indicating an amorphous region after anneal

Evidently, Fe does not fully compensate the Si electrically and RBS plots also indicate a thick amorphous layer. The data therefore suggests that conduction seen in the wafer is through hopping between the Fe sites, due to its high density. An anneal is also performed at 725 °C for 15 minutes, as suggested in the literature [13]; the sheet resistance is seen to be 70 Ω/□. At the chosen implant conditions, it is difficult to fully chemically compensate the high  $N^{++}$  doping. This could be due to formation of FeP precipitates, or due to the solubility limit of Fe in InP. It been reported that FeP precipitates were formed in CBE grown Fe doped InP above an Fe concentration of  $2 \times 10^{19} \text{ cm}^{-3}$ , and the resistivity is seen to drop [21]. Some authors [22] have indeed resorted to using damage Iron implants for isolation. They obtain 5 MΩ/□ for a room temperature implant. However the amorphous nature of the implant renders it unsuitable for subsequent crystalline regrowth.

$^{56}\text{Fe}$  is a heavy element (compared to  $^{28}\text{Si}$ ) and therefore creates a lot of damage to the crystal lattice during the implant process. Shown in Fig. 3.16 is the number of vacancies produced for Fe implanted at 150 keV, due to collisions with ions and recoiling atoms. The damage density is calculated for a dose of  $5 \times 10^{14} \text{ ions/cm}^2$  is

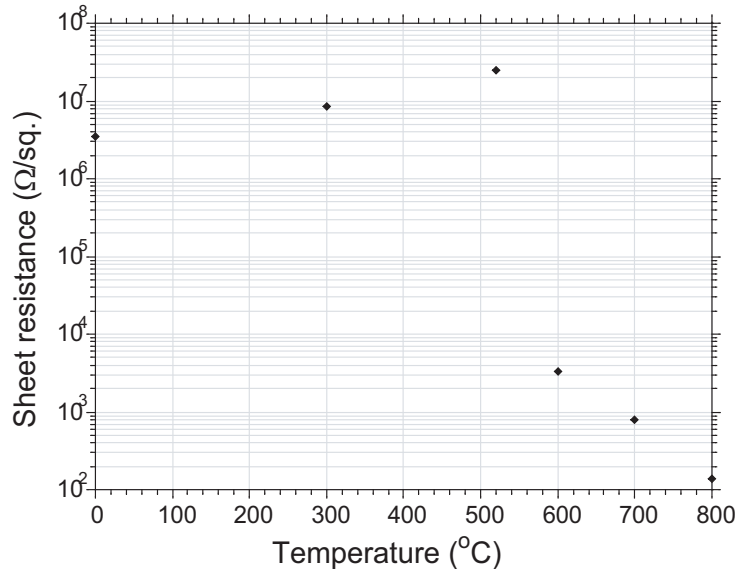


Figure 3.15: Resistivity with anneal for Fe implanted in 130nm  $N^-$ /80nm  $N^{++}$  InP at 190 keV/ $4 \times 10^{15}$  ions/cm<sup>2</sup>

to be  $\sim 5 \times 10^{22}$  atoms-cm<sup>-2</sup>, almost 100 % damage. This calculation ignores the dynamical annealing present during implants at elevated temperatures, but as seen from RBS and in Chapter 5, the implanted material is almost completely amorphous even when the implants are carried out at elevated temperature. This amorphous layer also influences certain chemical properties and this layer is seen to etch rapidly in BHF. It will be shown in Chapter 5 that annealing above 700 °C for long periods ( $\geq 5$  minutes) can anneal the damage completely for moderate doses (upto  $2 \times 10^{14}$  ions/cm<sup>2</sup>) and energies upto 150 keV. It was seen from the RBS plots and the resistivity data that it is difficult to anneal the damage when the implant dose is above a certain threshold. Previous work suggests that this dose increases with implant temperature [17].

One important parameter ignored in this discussion is the lateral straggle of Fe. At 190 KeV,  $\Delta R_z = 650 \text{ \AA}$ . The straggle becomes important when selectively masked implants are performed but throughout this work, Fe is implanted non selectively. Therefore, the lateral straggle is ignored here.

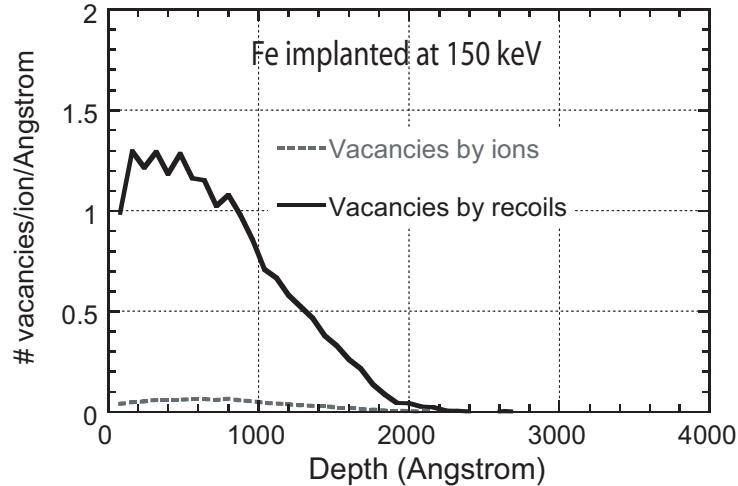


Figure 3.16: Damage in InP due to Fe implanted at 150 keV

### 3.1.4 Co-Implantation of Si and Fe

Si and Fe can be co-implanted as is done in Chapter 5. Since Si and Fe are activated only when they preferentially occupy group III sites, they both compete for In sites, when co-implanted. If Fe is implanted first, annealed and then Si is implanted, it has fewer group III sites to occupy. Hence the activation is reduced, albeit marginally. Co-implantation has yielded Schottky- $N^{++}$  diodes. The Si is implanted deep at 350 keV and Fe compensates the top 150 nm to form a SI region.

## 3.2 Mask for Implantation

Traditionally in the Si VLSI industry, photoresist (PR) is used to mask the implant. Since all the implants performed here are heated implants, it rules out any PR implant mask. Metal can be used as an efficient mask for implants. However, metals can be difficult to remove.  $SiO_2$  or  $Si_xN_y$  can also be used as a mask. However,  $SiO_2$  has a lower density than  $Si_xN_y$  and is therefore less efficient in stopping the ions. Thermally deposited  $Si_xN_y$  has a density of  $\sim 3.2$  g/cc, while plasma deposited  $Si_xN_y$  (at  $< 300^\circ\text{C}$ ) has a large concentration of hydrogen (10-35%) which lowers its density to 2.4 - 2.8 g/cc [23]. Here,  $Si_xN_y$  is used as an implant mask whenever selective implantation is carried out. Fig. 3.17 shows the profile of Si implants ( $300$  keV/ $6 \times 10^{14}$  ions/cm<sup>2</sup>) with  $1\mu\text{m}$   $Si_xN_y$  on InP. As seen from the figure, the density of Si at the  $Si_xN_y$ -InP boundary is  $\sim 1 \times 10^{16}$  cm<sup>-3</sup> which is not sufficient to cause N-

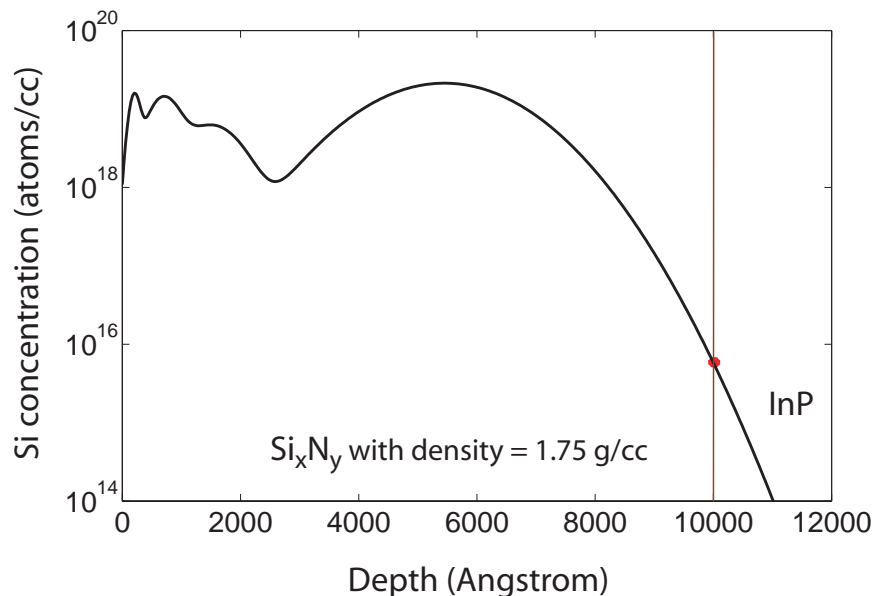


Figure 3.17:  $Si_xN_y$  implant mask for 300 keV Si implant in SI InP

type doping. Since the density of  $Si_xN_y$  from the PECVD is not exactly known, a much thicker mask than is predicted by *TRIM* is often used.

### 3.3 Growth on Implanted substrates

For fully crystalline growth, one of the requirements is a crystalline semiconductor and excellent surface morphology. As discussed in §3.1.1, the  $Si_xN_y$  has to be completely removed after anneal. Any Nitride remaining will cause poor growths and the Reflection-high energy electron diffraction (RHEED) signal patterns to be spotty. Fig. 3.5 is evidence that the process developed here results in complete removal of the  $Si_xN_y$  encapsulant after anneal. As will be seen in Chapter 4 and Chapter 5, anneal conditions developed for Si and Fe implants result in a crystalline semiconductor and in excellent growth morphology. Several published results exist for MBE growths on implanted InP substrates and epitaxial layers. Dodabalpur et al. demonstrated InAlAs/InGaAs HBTs with good DC characteristics grown on Si implanted InP substrates [24]. High speed HBTs and IC technologies have been demonstrated for layers overgrown on Si implanted epitaxial material [25, 26].

MBE growth is initiated after a surface treatment consisting of a 10 min. ultra



### CHAPTER 3. ION IMPLANTATION IN INP

violet(uv) ozone treatment followed by 5 min in BHF and 5 min. in DI water. As explained in §3.1.1, uv-ozone is an effective way of cleaning a surface of organic and non-organic materials and to form a stoichiometric oxide film. The BHF treatment removes this oxide. Prior to MBE growth standard oxide desorption is performed to rid the surface of remaining oxides. The RHEED patterns are seen to be streaky, indicative of excellent crystalline growth and the surface is observed under high resolution optical and electron microscopes to be comparable to fully epitaxial layers.

Having set a background into Si and Fe implantation in InP, Chapter ?? and Chapter 5 will explore practical applications of these implants for high speed devices.

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# 4

## Implanted Subcollector DHBTs

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HETROJUNCTION bipolar transistors are mostly fabricated in a triple mesa process. While these have achieved tremendous performance [1, 2] at the current scaling generation, there exists several issues that must be solved before scaling them further. Fig. 4.1 is an micrograph of a standard triple mesa HBT at UCSB [3]. In this device there is a significant portion of the base metal that lies outside the device active region. On the base access pad, is the base post which is  $\approx 1 \mu\text{m}$  in height. This connects the base metal to the top level metal interconnect. The base post, and hence the access pad have to be  $\geq 2 \mu\text{m}^2$  in order to achieve a reliable and low resistance contact with the metal interconnect. As shown in the cross sectional view of the device in Fig. 4.2, the base pad contributes to the base collector capacitance.

As discussed in Chapter 2, this base pad capacitance becomes an important parasitic as transistors are scaled. Low power logic can be achieved by using low current levels. However, to not impact the bandwidth, high current densities should be maintained. This necessitates the use of shorter length devices. Eqn.3.3 is repeated here.

CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTs

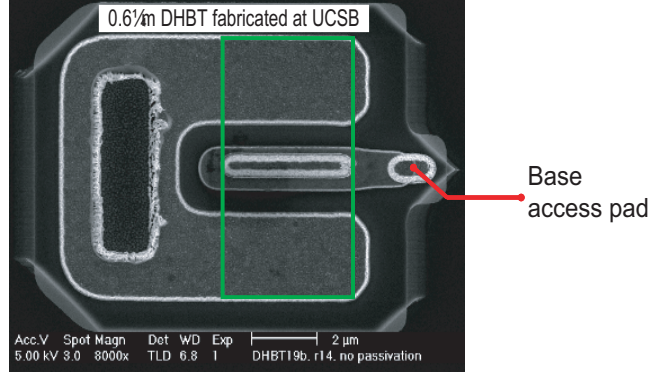


Figure 4.1: Planar view of a standard mesa HBT fabricated at UCSB

A major delay component determining digital logic speed is,

$$\hat{C}_{cb} \cdot \frac{W_{bm}}{W_{je}} \cdot \frac{\Delta V_{logic}}{J_e} + C_{cb,pad} \cdot \frac{\Delta V_{logic}}{J_e L_{je} W_{je}} \quad (4.0.0)$$

where  $\hat{C}_{cb}$  is the device capacitance per unit area (not including  $C_{cb,pad}$ , the access pad capacitance). For reasons explained, this base pad cannot be scaled. As seen from Eqn.3.3, as the length is decreased,  $C_{cb,pad}$  becomes a significant fraction of the total  $C_{cb}$ .

As detailed in Chapter 2, eliminating the pad capacitance reduces power consumption while maintaining the same bandwidth. Current state of the art mesa DHBTs have resulted in static frequency dividers operating at 150 GHz but consuming 600 mW [4]. In order to design a divider to operate at 150 GHz but at half the power, current mode logic (CML) can be employed for the data level and emitter coupled logic (ECL) for the clock level. With current mode logic, the transistors in the data level has a collector base swing of  $-\Delta V_{logic}$  to  $+\Delta V_{logic}$  as opposed to  $-\Delta V_{logic} + V_{BE}$  to  $+\Delta V_{logic} + V_{BE}$  for levels using emitter coupled logic. CML operation has greatly reduced maximum current density. From Eqn.3.3,  $C_{cb}/I_c$  increases and hence the delay. In order to maintain  $C_{cb}/I_c$ , a simple calculation and simulations indicate that  $C_{cb}$  has to be decreased by  $\sim 30\%$ .

There are several methods reported for eliminating this base access pad capacitance through the use of micro air-bridges [5, 6]. Fig. 4.2 shows a secondary electron micrograph of such as base pad. Typically in such processes, the subcollector under the feed lines from the base pad is removed by anisotropic wet etching. There are several disadvantages:

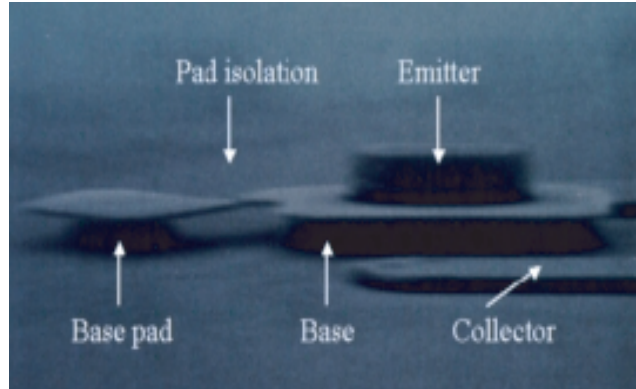


Figure 4.2: Air bridge to isolate the base access pad: Courtesy NTT, Japan

- The base feed lines are designed to be  $\approx 0.6-0.8 \mu\text{m}$  to ensure complete removal of the semiconductor below. These present a very large access resistance.
- The use of thin feed lines and the presence of air gaps, presents a reliability issue.

This technique is thus unsuitable for high yield, manufacturable processes.

An alternate idea is to isolate the base pad by selective implantation. Ion implantation is a viable method to achieve isolation in the active regions and has been successfully employed in GaAs-AlGaAs HBTs [7]. As discussed in Chapter 3, Fe implantation can be used to compensate the N-type collector and subcollector layers, where isolation is required Fig. 4.1. As detailed in the previous section a very high dose and energy of Iron is required to isolate  $N^{++}$  layers. Such Fe implants induces large defect density and is unsuitable where crystalline growth is required.

One of the main ideas explored in this thesis is the use of selective Si implants to eliminate the base pad capacitance.

## 4.1 Implanted subcollector HBT process

The process flow of the basic, implanted subcollector HBT is shown in Fig. 4.3. Starting with a template of Semi-Insulating InP, alignment targets for stepper based lithography are defined.  $Si_xN_y$  is deposited and patterns are defined by dry etch so that SI InP substrate is selectively implanted with Silicon(Si), an N-type dopant (Reasons for choosing Si are detailed in Chapter 3). Following the implant, the

## CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTS

$Si_xN_y$  implant mask is removed in BHF. The Si dopant is activated and the implant damage is removed, by a high temperature anneal in the RTA. In order that the Phosphorus not dissociate during anneal, the top and bottom of the wafer is covered with 40 nm  $Si_xN_y$ . This ensures that a highly doped, isolated subcollector region is formed only where devices are to be formed. In Fig. 4.1, Si is implanted inside the drawn boundary. The base access pad lies outside the subcollector implant boundary. This means that the subcollector is not present under this pad and there is no terminating plate of charge for the electric field lines from the  $p^{++}$  base. The capacitance due to the base pad is therefore nearly eliminated. The HBT drift collector, base, and emitter layers are then grown by Molecular Beam Epitaxy (MBE). Devices are fabricated in a standard, all wet etch mesa process. However with the selectively implanted subcollector, HBT isolation does not require a mesa etch. As seen from Fig. 4.3, this reduces the HBT mesa height by  $\sim 500$  nm significantly improving the planarity of the device and hence device yield in a complex IC process.

CHAPTER 4. IMPLANTED SUBCOLLECTOR HBTs

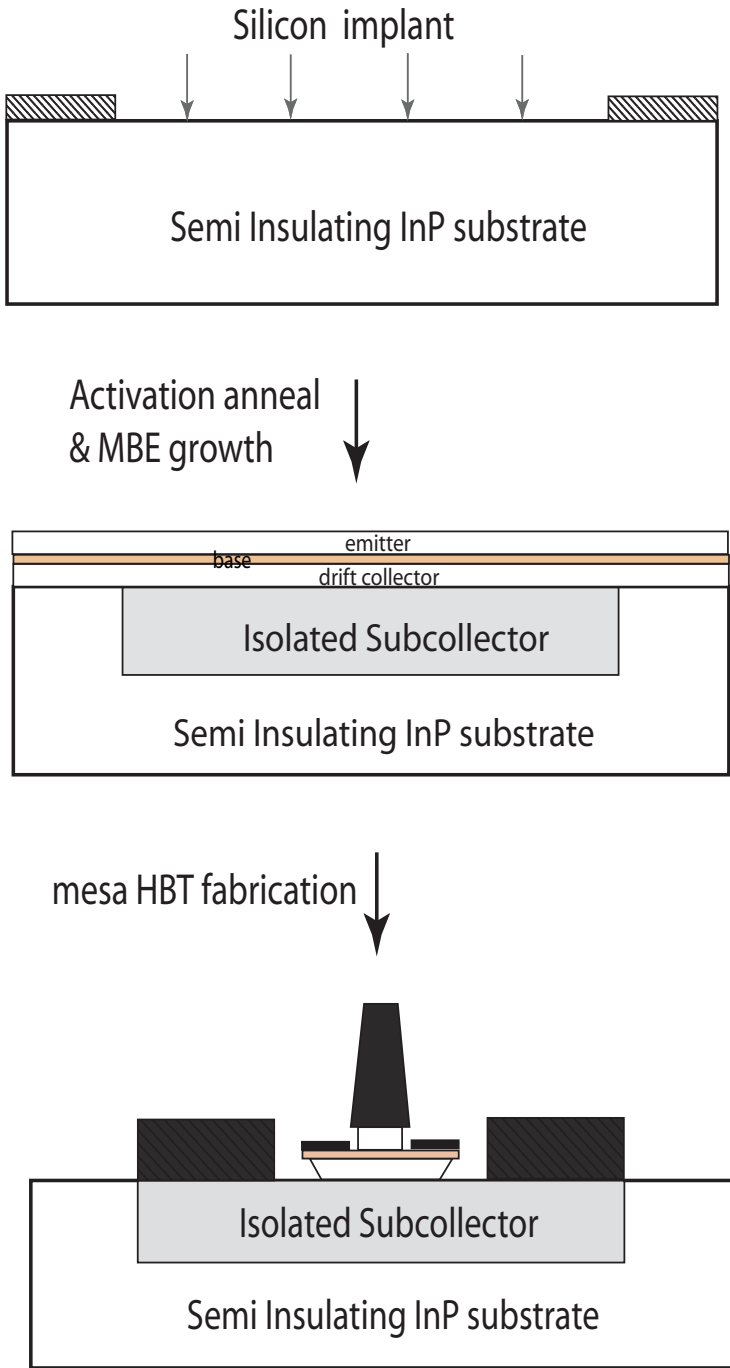


Figure 4.3: Basic implanted subcollector HBT process flow



## CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTS

Several criteria need to be satisfied for the implanted subcollector process.

- low sheet resistance subcollector
- high doping at the surface for low collector contact resistance
- crystalline, defect free semiconductor before growth
- excellent surface morphology comparable to InP substrate prior to growth
- planar surface with no recesses due to dry etches/implants/other processing
- high temperature stable alignment targets
- low contact resistivity ohmics to  $N^{++}$  InP
- good isolation between the  $N^{++}$  subcollectors
- defect free MBE growth  $\Rightarrow$  low  $I_{CBO}$ , DC characteristics similar to epitaxially grown HBTs

### 4.1.1 Si subcollector implant

From the expression for collector resistance in 2.3, the subcollector has to have high doping of  $\sim 1-2 \times 10^{19}$ , and a low sheet resistance comparable to a mesa DHBT  $\sim 10 \Omega/\square$ . Furthermore the subcollector needs to be isolated with very low inter device leakage  $\sim 1 \text{ pA}/\mu\text{m}$ . The requirements for the implant are thus summarized below.

- Low  $R_c \Rightarrow$  high doping  $\Rightarrow$  high implant fluence
- Low  $R_{sh} \Rightarrow$  thick subcollector  $\Rightarrow$  high implant energy
- Good isolation  $\Rightarrow$  excellent implant mask  $\Rightarrow$  thick  $Si_xN_y$

As discussed in Chapter 3 , the anneal should repair the crystal damage and also activate most of the dopants so that Si preferentially occupies the group III(In) site.

From the data in Table 3.2, the lowest sheet resistance when the Energy = 140 KeV (all other conditions are identical) is  $\sim 45 \Omega/\square$ . This is high compared to  $15 \Omega/\square$  for epitaxially grown subcollector in a standard mesa DHBT. In an effort to decrease the sheet resistance, the maximum implant energy is increase to 200 KeV to increase the implant depth. Following *TRIM* simulations, the distribution of the implanted Si ions in InP is shown in Fig. 4.4 for the implant conditions given in Table 4.1.

This is a multiple energy-multiple fluence implant.

CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTS

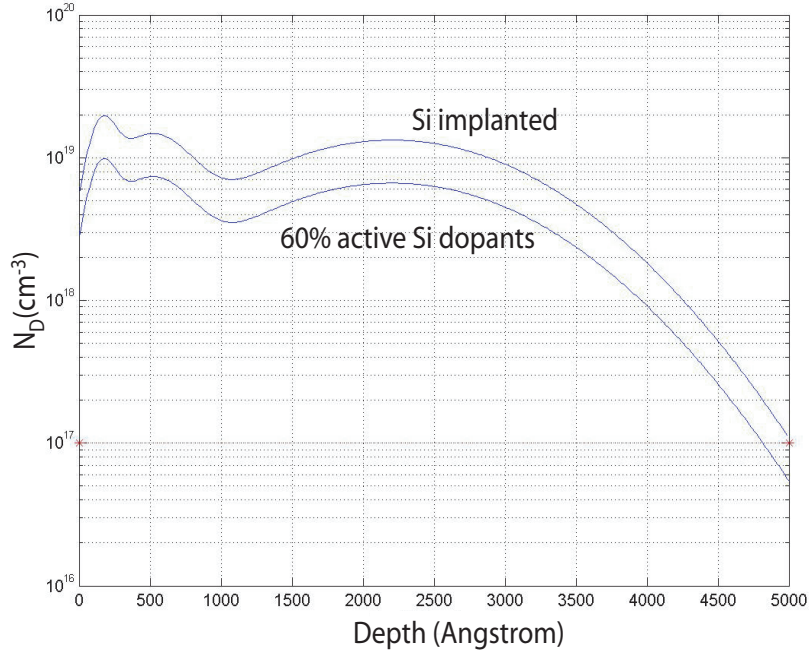


Figure 4.4: Concentration profile of Si in InP for implant conditions in Table 4.1

- The fluences and energies are chosen so as to obtain an overall concentration of  $\geq 2 \times 10^{19}$  over  $\sim 250$  nm. For epitaxially grown InP, the sheet resistance of such a layer would be  $\sim 15 \Omega/\square$ . The low energy component of the implant to ensure high doping at the surface for a good ohmic contact. This energy also ensures that the thickness of the  $Si_xN_y$  implant mask is not excessive ( $\approx 1 \mu\text{m}$ ). The implant fluences are kept within the known amorphization limits of Si implants in InP [8].
- All implants are done at an offset angle of  $7^\circ$  to prevent channeling through the substrate.
- During these implants the substrate temperature is maintained at  $200^\circ\text{C}$  to minimize the crystal damage due to implant.
- During these implants the current is always kept below  $10 \mu\text{A}$ , so that the substrate does not suffer from additional heating effects.

At this time, the highest known commercially available energy for Si implant at  $200^\circ\text{C}$  is 350 KeV. In order to simplify the design of the implant mask and the process,

## CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTS

Table 4.1: Implant conditions for basic, implanted subcollector HBT process

Implant species	Implant energy	Implant fluence	Offset angle	Implant temperature
Si	200 KeV	$3 \times 10^{14}$ ions/cm <sup>2</sup>	7 °	200 ° C
	40 KeV	$8 \times 10^{13}$ ions/cm <sup>2</sup>		
	10 KeV	$3 \times 10^{13}$ ions/cm <sup>2</sup>		
$I_{ions} \leq 10 \mu A$				

the above conditions are chosen for first generation transistors in this process. This is shown to yield sufficiently low values of  $R_{sh}$ . Extensive discussion on implants in InP etc. has already been provided in Chapter 3

### Sheet resistance

After this implant, a high temperature anneal is performed and the sheet concentration( $n_{sh}$ ), mobility( $\mu_n$ ), activation(%) and sheet resistance of samples, for the above implant conditions, are determined from standard four point Hall measurements and shown in Table 4.7. 800 °C for 30 seconds was chosen as the activation anneal for all Si im-

Table 4.2: Hall measurements of resistivity and mobility for Si implants

Anneal temperature (°C)	Anneal time (s)	Hall mobility ( $cm^2/V \cdot s$ )	Carrier density ( $cm^{-2}$ )	Sheet resistance ( $\Omega/\square$ )
800	30	722	$3.3 \times 10^{14}$	26
850	15	900	$3.1 \times 10^{14}$	22

plants. The activation is  $\approx 60$  %. The mobility is lower, and sheet resistance higher, when compared to epitaxially grown InP ( $\sim 1000$   $cm^2/V \cdot s$ ). Calculations of collector resistance showed that it is still dominated by the contact resistivity. Hence these implant and anneal conditions were chosen as a starting point. The anneal at 850 °C for 15 seconds, yielded a lower sheet resistance but the quality of the  $Si_xN_y$  cap was compromised giving rise to several anneal defects as seen in Chapter 3 . The sheet resistance obtained from Hall measurements correlates with sheet resistances measured using the transmission line method (TLM).

An anneal at 800 °C for 30s gives a resistance of  $\approx 25$   $\Omega/\square$ . It will be seen in Chapter 5 that the defects from the various dry etches and Si implant are sufficiently annealed out for these conditions. After implant and anneal, the active device layers are grown. There is no thick buffer as in a standard mesa DHBT process. From Atomic Force Microscopy(AFM) surface scans, the mean surface roughness for various anneal conditions. After annealing at 800 °C for 30s, the mean roughness is 1.8

## CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTS

nm. Virgin SI InP has a mean roughness of  $\sim 1.4$  nm. AFM scans are also performed to evaluate the uniformity of the surface after implant.  $Si_xN_y$  is the implant masked and is dry etched to open up the regions where Si is to be implanted, as seen in Fig. 4.3. The implanted surface is subjected to ion bombardment during dry etching and implantation. It is critical that the surface is fairly planar before growth since MBE involves line of sight deposition and hence growth in trenches is not straightforward. Fig. 4.5 shows a scan of such a selectively implanted section. The scan indicates rms planarity of  $\sim 5$ nm.

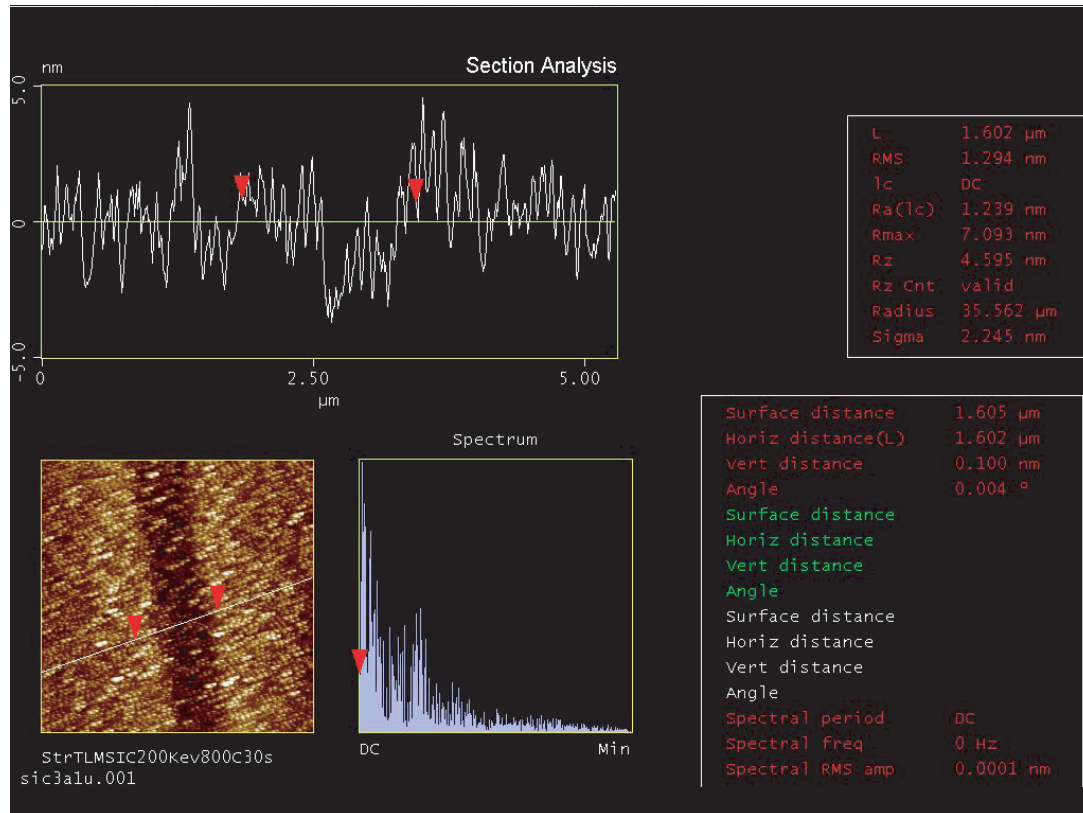


Figure 4.5: AFM scans of selectively implanted sections

### Device Isolation

$1\mu\text{m}$  thick  $\text{Si}_x\text{N}_y$  is the implant mask and from Fig. ?? in Chapter 3, this is seen to be sufficient to stop Si at the chosen conditions. Higher energy  $\Rightarrow$  thicker  $\text{Si}_x\text{N}_y$

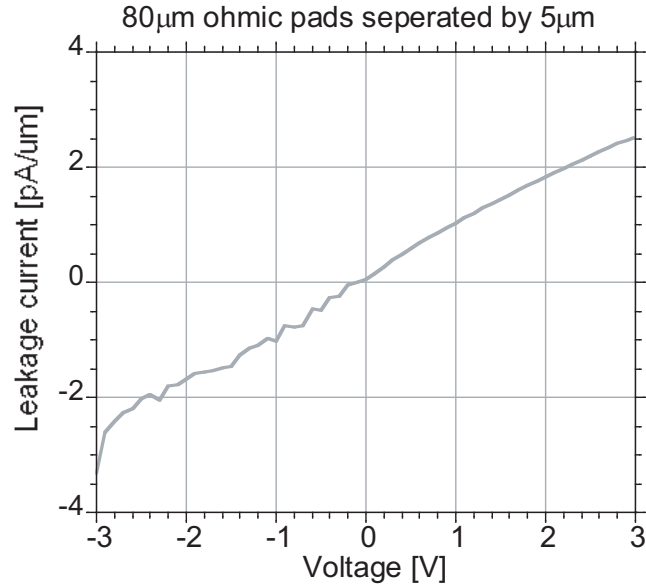


Figure 4.6: Device isolation between ohmic pads separated by  $5\mu\text{m}$

implant mask  $\Rightarrow$  unknown photoresist recipes to dry etch thick  $\text{Si}_x\text{N}_y$  in the reactive ion etcher (RIE) available at UCSB.

TLM patterns are formed to test the isolation of these subcollectors and hence the effectiveness of the  $\text{Si}_x\text{N}_y$  implant mask. Si is implanted under the  $N^{++}$  ohmic pads. Following activation by high temperature anneal, ohmic contacts are formed to InP. From Fig. 4.6, the leakage current in the regions masked during implant is  $\sim \text{pA}/\mu\text{m}$  at 3V, where the ohmic pads were separated by  $5\mu\text{m}$ .

### 4.1.2 Collector ohmic contacts

In a standard mesa DHBT process, collector ohmics are formed to  $N^{++}$  6.5 nm  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  to obtain low contact resistivities of  $\sim 10\Omega\cdot\mu\text{m}^2$ . As will be detailed in the next section, in the implanted subcollector DHBTs, the  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  layer is undoped and merely used as an etch stop layer. Collector contacts are made to  $N^{++}$  InP formed by the Si implant. Collector resistance is of a mesa DHBT with a

## CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTS

two sided collector is given by Eqn.2.3 in Chapter 2 ,

$$R_c = R_{c,contact} + R_{c,gap} + R_{c,spread} \quad (4.1.0)$$

$$R_c = \frac{1}{2} \frac{\sqrt{\rho_c \cdot R_{sh}}}{L_e}$$

where  $\rho_c$  ( $\Omega \cdot \mu\text{m}^2$ ) and  $R_{sh,s}$  ( $\Omega/\square$ ) are the specific contact resistivity and sheet resistance of the sub-collector. With  $R_{sh,s} \sim 25 \Omega/\square$ , base-collector spacing of  $0.5 \mu\text{m}$ , base mesa width of  $1.35 \mu\text{m}$ , contact resistivity  $\geq 10 \Omega \cdot \mu\text{m}^2$ , and assuming that the collector contact width is  $\gg L_T$  the ohmic transfer length, the various components of  $R_c$  are given by

$$R_{c,contact} \geq 8\Omega \cdot \mu\text{m}, R_{c,gap} \approx 6\Omega \cdot \mu\text{m}, R_{c,spread} \approx 2.8\Omega \cdot \mu\text{m} \quad (4.1.0)$$

The contact resistance dominates even for contact resistivities as low as  $10 \Omega \cdot \mu\text{m}^2$ . Since collector contacts are made to wide bandgap InP, contact resistance assumes significance.

The effect of collector contact resistance on digital logic speed is given by,

$$f_{CLK}^{-1} \propto \dots + C_{cb} \cdot \left( \frac{\Delta V_L}{I_c} + R_{c,contact} \right) \quad (4.1.0)$$

At the current scaling generation [4], for  $\Delta V_{logic} = 300 \text{ mV}$  at an operating current of  $I_c = 5 \text{ mA}/\mu\text{m}$ , the load resistance  $R_L = \Delta V_L/I_c$  is  $40 \Omega \cdot \mu\text{m}$ . The collector contact resistance is in series with this load resistance. For the above values,  $R_{c,contact}$  constitutes  $\sim 16 \%$  of the delay term given by Eqn.4.1.2. Similarly,  $C_{cb} \cdot R_{c,contact}$  delays reduce  $f_\tau$  and  $f_{max}$ .

Collector ohmics are made to  $N^{++}$  InP. Since the barrier height are larger for wideband gap semiconductors, alloyed ohmics(AuGe) are typically employed to contact wide bandgap semiconductors.  $Au_{0.88}Ge_{0.12}$  forms a eutectic at  $365 \text{ }^\circ\text{C}$ . In GaAs, the Au reacts with GaAs leaving behind a large concentration of Ga vacancies and Ge diffuses, occupies the Ga sites and heavily dopes the underlying semiconductor N-type[9]. A thin interfacial Ni(1-5 nm) is used as the wetting layer and also serves to enhance the diffusion of the GeAu alloy[10]. However, the exact mechanism of ohmic formation with alloyed contacts is not well understood in N-InP.

Non alloyed ohmics such as TiPtAu to N-type InP typically yield high contact resistivities  $\sim 800 \Omega \cdot \mu\text{m}^2$  [11]. Sputter cleaning, with an inert gas, the surface before contact metal deposition reduces the contact resistivity of such non alloyed ohmics to  $\sim 40 \Omega \cdot \mu\text{m}^2$  [12]. This is attributed to the formation of In rich surface that is degenerately  $N^{++}$  doped. In contrast, alloyed GeAu ohmics yield low contact

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Table 4.3: Contact resistivity of various metalization schemes to Si implanted InP

S.No	Metalization	Anneal (°C)	$R_{sh}$ ( $\Omega/\square$ )	$\rho_c$ ( $\Omega \cdot \mu\text{m}^2$ )
1	Ni(20nm)/Ge(50nm)/Au(50nm)/Ni(20nm)/Au	300	23	90-100
2	Ni(5nm)/Ge <sub>0.12</sub> Au(~80nm)/Ni(20nm)/Au	not annealed	16	200-230
		320	16	45-50
		360	16	25-30
		400	16	12-18
3	Pd(3nm)/Ti(17nm)/Pd(~17nm)/Au	300	10	75-100
4	Ge(1nm)/Ni(5nm)/Ge(50nm)/Au(50nm)/Ni(20nm)/Au			
5				
6				
7				
1. Si implanted and annealed with the conditions given in subsection ?? 2,4. Si implanted at a higher energy $\sim 325$ KeV and annealed so that $R_{sh} \sim 15 \Omega/\square$ 3. Epitaxially grown $N^{++}$ InP subcollector, with $R_{sh} \sim 10 \Omega/\square$				

resistivities to Si implanted InP of  $\sim 7 - 10 \Omega \cdot \mu\text{m}^2$ . The low energy component(10 keV) of the Si implant employed here (see previous section) ensures a high surface doping for low contact resistance. Since collector contact formation to InP is not trivial, several experiments are carried out and the results are summarized in Table 4.3.

It is seen from the above data that the lowest contact resistivity is obtained with the AuGe eutectic annealed at 400 °C. However, it is seen that such anneals of the collector ohmics results in an increase in the base and emitter resistivities and is shown in Table 4.4. Further for alloyed ohmics, high anneals result in pits and poor surface morphology of the metal contacts.

Annealing at or above 360 °C causes the base contact resistivity to substantially increase. The base is doping graded from  $8 \times 10^{19}$  to  $5 \times 10^{19}$  from the emitter side to the collector. As the anneal temperature increases, the thin interfacial Pd diffuses, and forms contact with lesser doped material. The Pd and Ti can diffuse through the thin base at sufficiently high anneals. Shown in Fig. 4.7 are plots of the base collector leakage current showing the effect of the various anneals. At 400 °C, the base metal nearly shorts the collector. GeAu contacts require anneals above the eutectic temperature to obtain low contact resistivity. As seen from the above table,

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Table 4.4: Contact resistivity of base, emitter and collector ohmics at various anneals

Anneal °C	Emitter $\rho_c$ ( $\Omega \cdot \mu\text{m}^2$ )	Base $\rho_c$ ( $\Omega \cdot \mu\text{m}^2$ )	Collector $\rho_c$ ( $\Omega \cdot \mu\text{m}^2$ )
	Ti(20nm)/Pd(40nm)/Au	Pd(3nm)/Ti(17nm)/Pd(17nm)/Au	Ni(5nm)/Ge <sub>0.12</sub> Au(~80nm)/Ni(20nm)/Au
unannealed	5	12	200-230
320	5	18	45-50
360	5	55	25-30
400	50	270	12-18

Ni/Ge/Au where ge and Au are deposited as separately yields the lowest resistivity at 300 °C. For the initial process runs, these were chosen as collector contacts. This is similar to regular non-alloyed contacts. In order to use the GeAu eutectic, the base and collector ohmic processes need to be revised. Either the base ohmic must be redesigned to withstand high temperature anneals or new collector ohmics must be developed to  $N^{++}$  InP that yield very low contact resistivities without the need for an anneal.

### 4.1.3 Alignment targets

All features are defined by lithography using the i-line (365nm) GCA wafer stepper in UCSB nanofabrication facility. As HBT feature sizes are scaled, alignment becomes critical. With the Dark field alignment system(DFAS) on the GCA stepper, alignment error better than 0.15  $\mu\text{m}$  is achieved. This system delivers the best alignment, when there is sufficient contrast between the targets and the field. This requires the targets to be at a significantly different height from the field. The first step of the implanted subcollector HBT process, consists of creating alignment targets subsequently used to align the wafer so as to correctly define features for the subcollector implant. The alignment targets must withstand the dry etch, high temperature (800 °C) processing steps and MBE regrowth and must have sufficient contrast. Etching into the substrate is the most straightforward method of forming alignment targets. However, the fear that subsequent growth might reduce its step height and affect its contrast precluded its usage. The  $\geq 0.7 \mu\text{m}$  emitter metal also defines targets in a standard mesa process and provides excellent contrast for precise local alignment. However, most metal targets, such as Ti or Au are not compatible where a high temperature anneal is required [13]. The refractory metals, Tungsten, Molybdenum and Tantalum seem to be excellent choices. Molybdenum is etched in BHF which is used to remove the  $Si_xN_y$  implant mask while Tantalum deposition is not straightforward



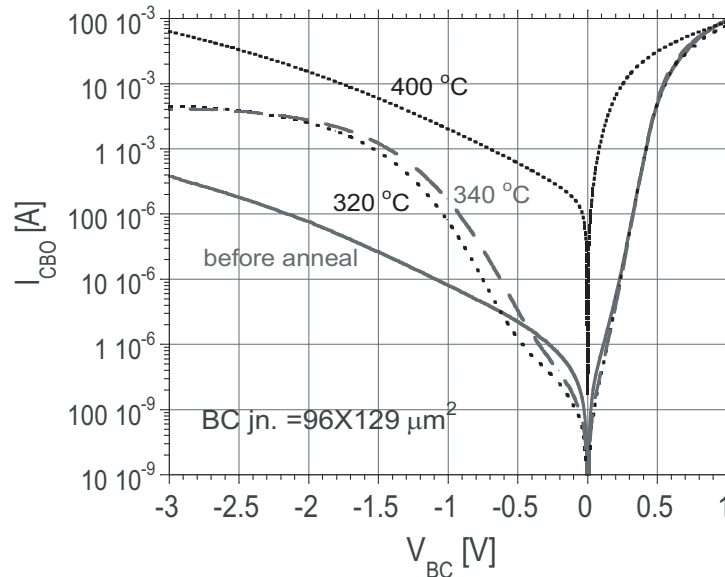


Figure 4.7: Base collector leakage upon annealing

and it is highly toxic. Hence the only viable option was Tungsten ( $W$ ).

Pure  $W$  is a hard refractory metal with the highest melting point (3422 °C) of all metals. It has very low chemical activity at room temperatures and is attacked slightly by most mineral acids. Owing to its high melting point and its refractory nature,  $W$  is used extensively in high-temperature applications. In the implanted subcollector process,  $W$  was not attacked by any of the acids or bases used. The first process runs were carried out using  $W$  where 0.3  $\mu\text{m}$  Tungsten is sputtered and dry etched to form the alignment targets. These targets however did not withstand the high temperature anneal step to activate Si, and formed defective outgrowths as shown in Fig. 4.8. Due to these outgrowths of metal, the DFAS local alignment system did not recognize the targets. This presented a huge problem since fine alignment of  $\leq 0.15 \mu\text{m}$  was required which was obtainable reliably only by using the DFAS. One option was to reduce the anneal temperature for Si activation but this meant a compromise to device performance. Studies and analysis of these  $W$  targets are carried out using energy dispersive x-ray analysis (EDX). Detecting the characteristic fluorescence x-rays emitted from the sample as a result of excitation by the imaging electron beam, elemental analysis using EDX is performed. Fig. 4.9 shows the EDX spectra of the overhang metal features on the targets after anneal. A huge Oxygen concentration is detected. It is found that residual stress and structural prop-

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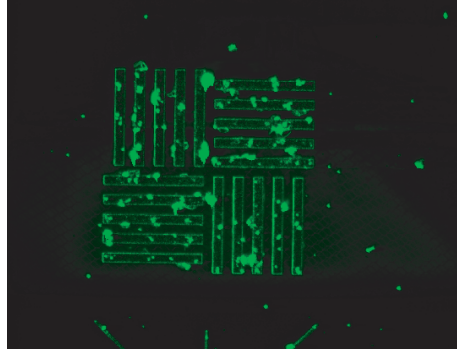


Figure 4.8: Tungsten alignment targets after high temperature anneal

erties of tungsten films, is a function of the sputtering gas pressure. There is also found to be a strong correlation between the stress and the microstructure of these films. The amorphous phase has been found to contain over 20% Oxygen and is reported to be metastable [?]. The thermal properties could thus be severely impacted by the presence of large oxygen impurities in the sputtered Tungsten.

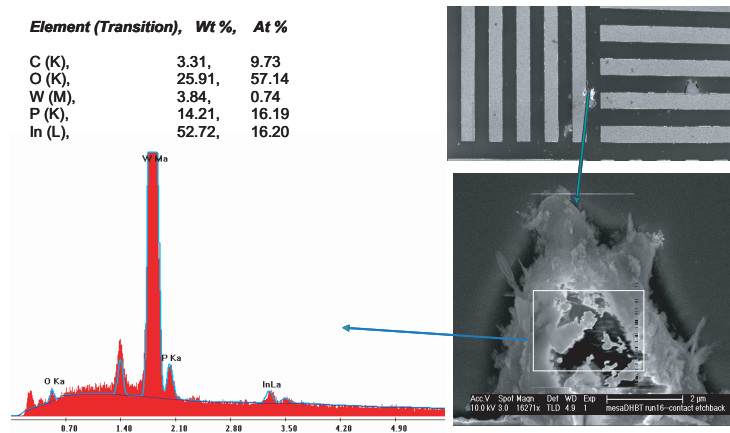


Figure 4.9: EDX spectra of the defects on the Tungsten targets after anneal

Ti(10%)W was reported to be stable after annealing at 800 °C [15]. TiW alloy has similar chemical properties to W. It could be easily dry etched and is not attacked by any of the chemicals used in this process. Furthermore, these targets had better contrast on the stepper's local alignment system than W, which is a dull grey metal.

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TiW was sputtered at low pressures to a height of  $\sim 0.3 \mu\text{m}$ . These targets withstood an annealing temperature of  $800^\circ\text{C}$ . They are stable (Fig. 4.10) after MBE growth, and are resolved by the stepper's alignment system.

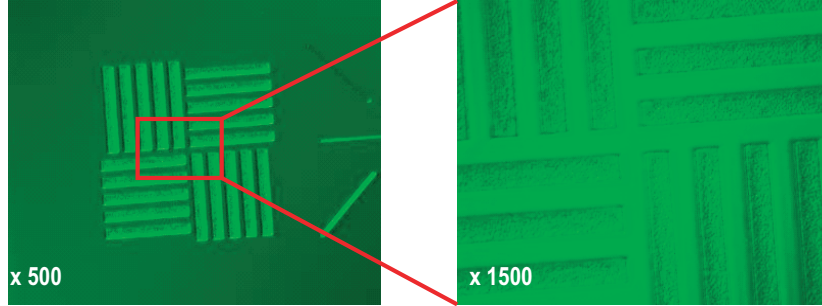


Figure 4.10: TiW alignment targets after high temperature anneal and MBE growth

### 4.1.4 Design of RF mask

The region where the base pad lies has to be masked during Si implant. Si has to be implanted under the collector contacts and in the active device region (Fig. ??). The Si implant also has to be present under the emitter junction. In actuality, it has to extend beyond the length of the emitter junction due to current spreading. The length of the subcollector mask is determined as follows.

$$L_{sc} = L_E + T_c - \Delta x_E - \Delta x_{straggle} + 2\Delta x_{align} \quad (4.1.0)$$

where  $L_E$  is the length of the emitter as on the photomask in Fig. 4.11,  $\Delta x_E$  is the longitudinal undercut of the emitter,  $\Delta x_{straggle}$  is the straggle due to implant,  $\Delta x_{align}$  is the alignment tolerance. Since the subcollector and emitter are individually referenced to separate zero level targets, the alignment error doubles. The implant straggle from TRIM plots is seen to be  $\sim 0.2 \mu\text{m}$ . With  $\Delta x_E = 0.4 \mu\text{m}$ ,  $\Delta x_{align} = 0.2 \mu\text{m}$ .

$$L_{sc} = L_E + T_c - 0.2 \quad (4.1.0)$$

To accommodate collector thicknesses below 200 nm, the length of the subcollector photomask is designed such that,  $L_{sc} = L_E$ . In order to compare the effectiveness of the subcollector implant process, the base pad is not isolated in some devices, so that the region under the base pad is not masked during implant.

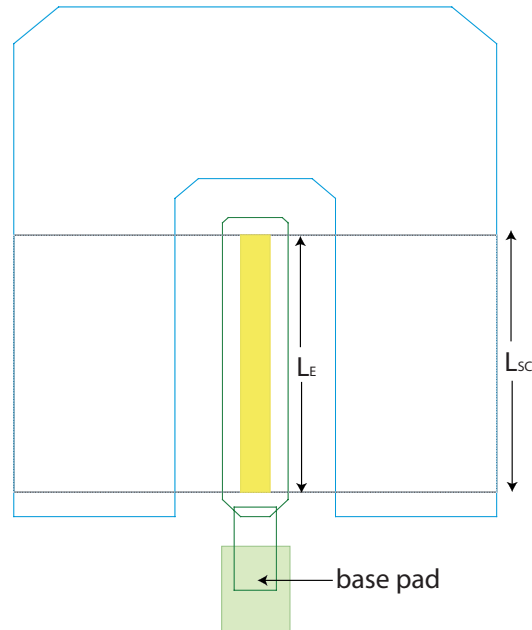


Figure 4.11: Layout of implant mask, seen with the emitter, base and base access pad

## 4.2 HBT design for implanted subcollector

The design of each layer in an HBT is detailed in Chapter 2 . With the lithographic capabilities available at the time  $0.6 \mu\text{m}$  features can be reliably processed at UCSB, even for complex ICs. The implanted subcollector DHBTs have significant performance advantages when devices are scaled. However, the first generation transistors are intended to compare the advantages of this process against the mesa technology. In order to make intelligent assessments, performance comparisons has to be made against existing transistor results. Fast HBTS have been reported at UCSB, with collector thickness  $T_C = 200\text{nm}$  [16],  $150 \text{ nm}$  [17],  $120 \text{ nm}$  [3],  $100 \text{ nm}$  [18].

Static frequency dividers with  $\Delta V_{logic} = 300 \text{ mV}$ , demonstrated a maximum clock speed of  $140 \text{ GHz}$ [4]. These have a  $C_{cb}/I_c$  of  $\approx 0.5 \text{ ps/V}$ . As stated earlier,  $C_{cb}$  reduction techniques results in superior bandwidth of digital logic circuits and the best performance metric for implanted subcollector DHBTs is maximum digital

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logic speed. The minimum voltage swing is given by,

$$\Delta V_{logic} \geq 2J_e \cdot \rho_e + 6 \frac{kT}{q} \quad (4.2.0)$$

A typical value of emitter contact resistivity repeatedly obtained is  $\rho_c \sim 10 \Omega \cdot \mu\text{m}^2$ . If  $J_e \sim 8 \text{ mA}/\mu\text{m}^2$ ,  $J_e \cdot \rho_c \sim 80 \text{ mV}$ . The device operating temperature is  $\sim 400\text{K}$ , therefore  $6kT/q \sim 200 \text{ meV}$ . The noise margin is only  $\sim 20 \text{ mV}$ . There for these values of contact resistivity, the maximum operating current density has to be maintained below  $8 \text{ mA}/\mu\text{m}^2$ . Furthermore the maximum operating power density of the ECL dividers is  $\sim 1.1\text{V} \times 8 \text{ mA}/\mu\text{m}^2 \approx 9 \text{ mW}/\mu\text{m}^2$  and such power densities should be supported in the layer designs.

As stated earlier, in order to compare against existing technology, the emitter and base layers for the first generation implanted subcollector DHBTs are identical to the mesa DHBTs reported [3]. The emitter base junction is abrupt, also due to ease of processing. (Selectively wet etching a  $\text{In}_{0.52}\text{Al}_{0.48}\text{As}$ - $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  grade is not straightforward). The base is designed to be 30 nm. For the contact metallurgy used and doping levels obtainable, this offers the best compromise between  $\tau_b$  and  $R_{bb}$ . The collector design was an open choice. However, the maximum operating current density has to be kept below  $8 \text{ mA}/\mu\text{m}^2$ . 120nm collectors have a maximum operating current threshold of  $\sim 8 \text{ mA}/\mu\text{m}^2$ . From the scaling laws, in order to maintain the highest current density and lowest  $C_{cb}$ , 120 nm is the preferred choice. The minimum resolvable lithographic limit with good device yield for ICs at the UCSB nanofab facility, at this time is  $\sim 0.6 \mu\text{m}$ . Minimum repeatable contact resistivities obtainable are,  $\rho_c \sim 10 \Omega \cdot \mu\text{m}^2$ . The lowest base resistivity for state of the art C doped, MBE grown  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  layer is  $18 \Omega \cdot \mu\text{m}$ . For 30 nm bases, this yields an ohmic contact length of  $\sim 0.13 \mu\text{m}$ . With an alignment tolerance of  $0.15 \mu\text{m}$ , the base ohmic has to be at least  $0.3 \mu\text{m}$  wide on either side of the emitter for low base resistance. For dividers, the emitter widths are designed to be  $\geq$  than the present lithographic of  $0.6 \mu\text{m}$ .

With new lithographic capabilities acquired recently at UCSB, and in conjunction with better specific contact resistivities that have been recently obtained, the implanted subcollector DHBTs will greatly benefit future laterally and vertically scaled transistors.

### 4.2.1 Design of InGaAs etch stop

In standard mesa DHBTs, the  $N^{++}$  subcollector is a composite structure and consists of highly doped  $N^{++}$  6.5 nm InGaAs over 300 nm InP. The  $N^{++}$  InGaAs serves as the contact layer and also as an etch stop to selectively stop on the subcollector during the second mesa etch.

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In the implanted subcollector DHBTs, the  $N^{++}$  InP is selectively implanted.  $N^{++}$  InGaAs cannot be present on these implanted subcollectors. This layer which is present and  $N^{++}$  doped, will form a terminating electrode under the base pad. This negates the entire idea of having selectively implanted  $N^{++}$  regions to eliminate the base pad capacitance. However an InGaAs layer is required as an etch stop. Therefore this InGaAs layer has to be necessarily undoped.

The InGaAs layer is to be sandwiched between InP layers and has a conduction band discontinuity of 0.25 eV with InP. A thin layer of InGaAs therefore forms a quantum well of electrons with a depth of 0.25 eV. It is critical that there be no electron/hole pile up in this quantum well. This means that the ground level for electrons in the quantum well must be much higher than the electron Fermi level. For a quantum well with infinite barriers, the ground state energy is given by

$$E_0 = \frac{\hbar^2 \pi^2}{2m_e^* L^2} \quad (4.2.0)$$

where  $L$  is the thickness of the quantum well. By reducing the thickness of this well, the ground state energy can be increased. But, for a finite well, it can be shown that there always exists at least one energy level. It can be shown that the number of levels [19] is given by,

$$N = 1 + \text{Int} \frac{2k_0 a}{\pi} \quad (4.2.0)$$

The ground state energy level for a quantum well of finite depth is given by,

$$E_0 = \frac{\hbar^2 k^2}{2m_e^*} \quad (4.2.0)$$

where the wavevector  $k$  is defined by

$$\frac{kL}{\cos kL} = k_0 L \quad (4.2.0)$$

and  $k_0$  is given by

$$k_0 = \sqrt{\frac{2MV_0}{\hbar^2}} \quad (4.2.0)$$

$V_0$  is the height of the well, here  $V_0 = \Delta E_c = 0.25 \text{ eV}$ . For example, in the extreme case of  $L = 0$ ,  $E_0 = V_0$ . For  $L = \infty$ ,  $E_0 = 0$ . Thus  $0 \leq E_0 \leq V_0$ ; there is at least one energy level in a quantum well with finite barriers. Simulations were carried out in 1-D Poisson to calculate the electron concentration in the InGaAs quantum well for quantum wells of width = 4nm and 5nm, and shown in Fig. 4.12. The InGaAs layer has to be thick enough to be an effective etchstop during the wet etching of the

#### *CHAPTER 4. IMPLANTED SUBCOLLECTOR DHBTS*

InP collector. The ground state energy level is 0.13eV above the Fermi level for an InGaAs layer of thickness 4 nm while the electron concentration is  $\sim 6 \times 10^{15}$  which is much lower than the collector doping of  $\sim 3 \times 10^{16}$ . The undoped InGaAs etch stop layer is designed to be 3.5 nm.

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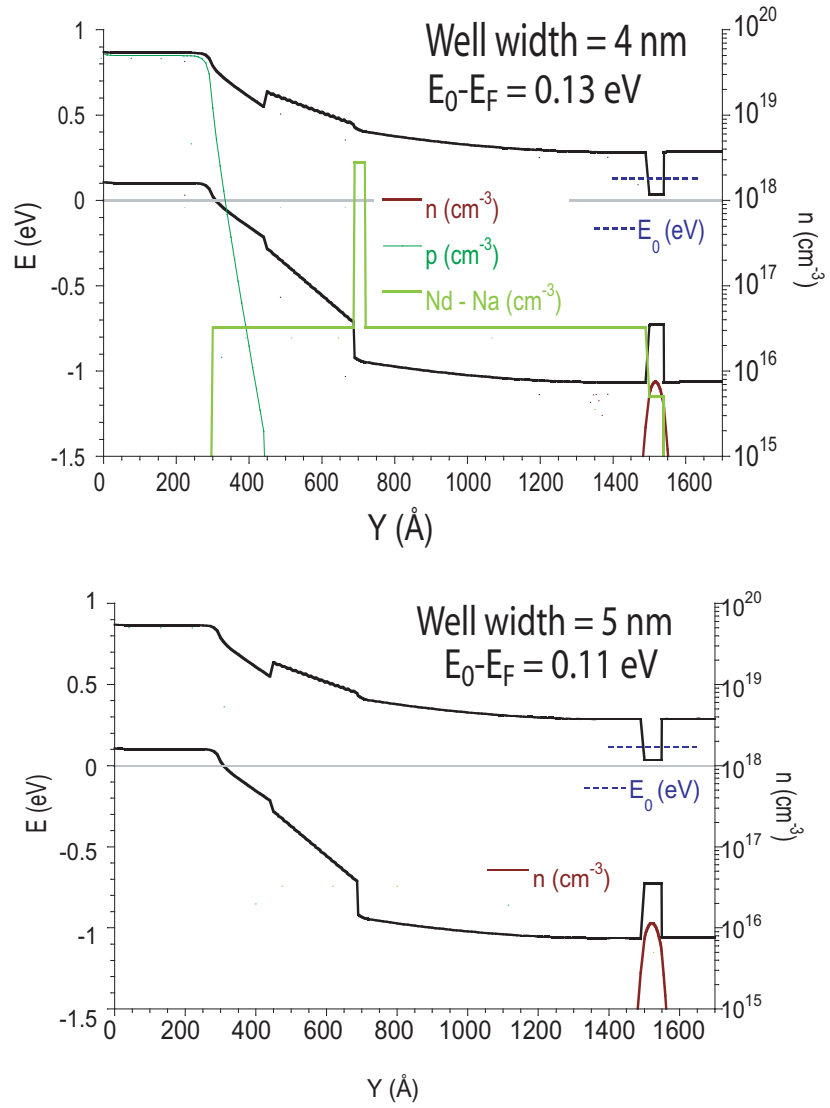


Figure 4.12: Electron concentration and Ground state energy level of quantum wells (4nm and 5nm) of UID InGaAs sandwiched between the collector and SI InP



### 4.3 Large area HBTs with implanted subcollector

In order to evaluate the layer structure, and growth on implanted substrates, Si InP is blanket implanted with Si at the conditions given Table 4.1. The wafer is annealed at 800 °C for 30 sec and subsequently growth was done by MBE at UCSB. The HBT layer structure consisted of a 150 nm collector and 30 nm base. Large area devices are fabricated where the emitter, base and collector are defined by 2 wet etches, followed by a single, common metalization (PdTiPdAu). From TLMs,

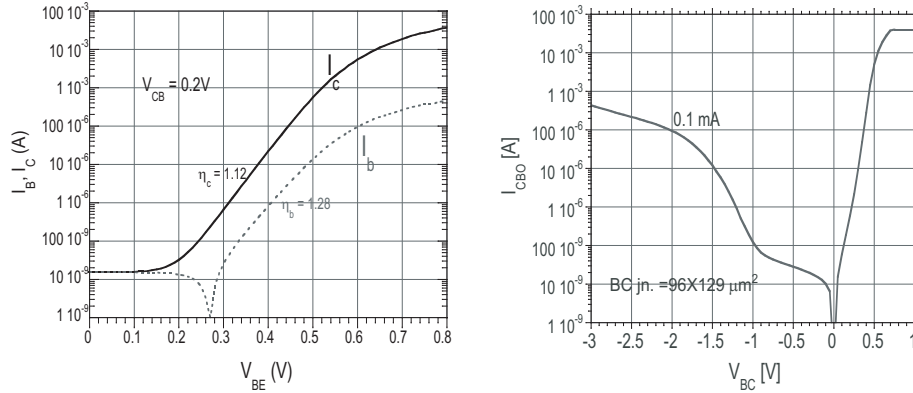


Figure 4.13: DC characteristics and base-collector leakage

the specific contact resistivities are  $5\text{-}10 \Omega \cdot \mu\text{m}^2$  while the base sheet resistance is very high  $\sim 3600 \Omega/\square$ . This is believed to be due to the surface depletion of  $\sim 10$  nm and chiefly due to miscalibration of the base Carbon doping and is corrected in subsequent growths. Fig. 4.13 shows the gummel characteristics of these large area HBTs. These devices show a gain of  $\approx 70$ . This high gain is further indicative of lower Auger recombination and hence low base doping. The plot on the right in Fig. 4.13 indicates that the leakage current at 2V is  $\sim 8 \times 10^{-6} \text{ mA}/\mu\text{m}^2$ . Extrapolating this for small area devices with a base collector area of  $1.3 \times 5.25 \mu\text{m}^2$ ,  $I_{CBO}$  is  $\sim 5.6 \times 10^{-5} \text{ mA}$  and  $I_{CEO}$  is  $\sim 3 \times 10^{-3} \text{ mA}$  which is a very low compared to operating currents of  $\geq 10 \text{ mA}$  and is hence indicative of a low base collector leakage and hence good growth on these implanted substrates. The very high base resistance is evident from the deviation from exponential characteristics even at moderate base emitter forward bias voltage of 0.8V.

Fig. 4.14 shows the Capacitance Voltage (CV) characteristics and the doping profile of the collector is extracted from the CV data. The data measured from the large area devices clearly illustrate that growth on implanted substrates is comparable to

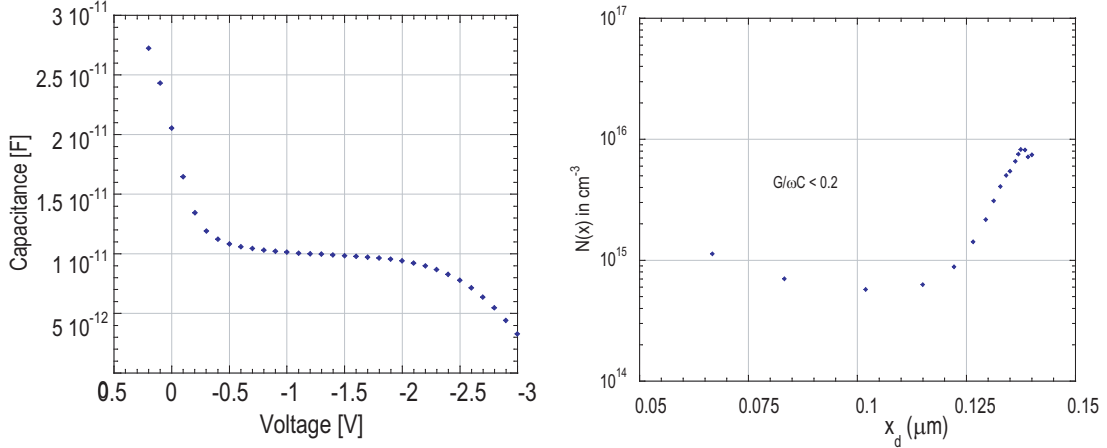


Figure 4.14: CV characteristics and doping profile

fully epitaxial structures.

## 4.4 RF process and device results

Large area devices have shown low leakage and good DC gain. Small area devices were processed with the layer structure shown in Table 4.5.

### 4.4.1 RF process

TiW alignment targets are defined, and following the process described in §4.1 and surface treatments elucidated in Chapter ??, growths were carried out at UCSB's MBE facility. A brief description of the process after growth is given below. The details are given in Appendix ?. The emitters are accurately aligned to the TiW alignment targets. Alignment targets are also defined with the emitter metal and all further alignments are referenced to these. The emitter layer is then wet etched down to the base.  $\sim 0.2 \mu\text{m}$  Base contacts and a base post of  $\sim 1 \mu\text{m}$  are deposited. Following this, the base mesa etch is performed down to the implanted InP subcollector. Collector contacts are defined using Ge/Ni/Ge/Au/Ni/Au. These were annealed only at a temperature of  $300^\circ\text{C}$  to not impact the base contact resistivity. After the collector post is deposited, the wafer is planarized with Benzo Cyclo Butene (BCB) which is etched down to barely expose the metal posts.  $100 \text{ nm}$  of  $\text{Si}_x\text{N}_y$  is deposited to improve Metal-1 (M1) adhesion and to form capacitors. The first level of intercon-

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nect M1 then contacts the devices after forming vias in  $SixNy$ . Devices are tested after this metalization step.

### 4.4.2 Device Results - Si Implanted subcollector DHBTs

The complete layer structure in the active device region are given in Table 4.5. The corresponding band diagram is depicted in Fig. 4.15. The projected  $J_{KirK} = 7 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0.0 \text{ V}$  and  $J_{KirK} = 11 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0.6 \text{ V}$ . The expected capacitance per unit area from Fig. ?? is  $\approx 1 \text{ fF}/\mu\text{m}^2$ .

Table 4.5: Layer structure: Si Implanted subcollector, 120 nm collector, 30 nm base

Thickness (nm)	Semiconductor Composition	Doping ( $\text{cm}^{-3}$ )	Description
5	$\text{In}_{0.85}\text{Ga}_{0.15}\text{As}$	$> 4 \cdot 10^{19}:\text{Si}$	Emitter cap
15	$\text{In}_x\text{Ga}_{1-x}\text{As}$	$4 \cdot 10^{19}:\text{Si}$	Emitter cap
20	$\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$	$4 \cdot 10^{19}:\text{Si}$	Emitter cap
80	InP	$3 \cdot 10^{19}:\text{Si}$	Emitter
10	InP	$8 \cdot 10^{17}:\text{Si}$	Emitter
40	InP	$5 \cdot 10^{17}:\text{Si}$	Emitter
30	InGaAs	$4 - 7 \cdot 10^{19}:\text{C}$	Base
15	$\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$	$3.25 \cdot 10^{16}:\text{Si}$	Setback
24	InGaAs / InAlAs	$3.25 \cdot 10^{16}:\text{Si}$	B-C grade
3	InP	$3 \cdot 10^{18}:\text{Si}$	Delta doping
78	InP	$3.25 \cdot 10^{16}:\text{Si}$	Collector
3.5	$\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$	undoped	Etch Stop
Substrate	Si implanted InP		Subcollector

The TLM measurements of the collector and, of the pinched and non-pinched base TLMs are given in Fig. ?. Isolation between ohmic pads separated by  $10 \mu\text{m}$  is  $\sim 20 \text{ pA}/\mu\text{m}$ .

Fig. 4.17 is a plot of the DC  $I_C - V_{CE}$  characteristics. There is no evidence of gain compression till  $\approx 9 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0\text{V}$ . The Gummel curves are plotted for  $V_{cb} = 0\text{V}$  and  $V_{cb} = 0.3\text{V}$  Fig. 4.18. The collector and base ideality factors are  $\eta_c = 1.15$  and  $\eta_b = 1.65$ . These are in line with those obtained for fully epitaxial DHBTs. The leakage current  $I_{CEO}$  from the gummel curves is  $\approx 90 \text{ pA}$  at  $V_{cb} = 0.3\text{V}$ . The DC current gain of the transistor  $= I_C/I_B$  from the gummel curves is shown in Fig. 4.19. The HBT's gain compression point is  $\approx 6 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0\text{V}$  and  $7 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0.3\text{V}$ .

Shown in Fig. 4.20 are  $I_{CEO}$  and  $I_{CBO}$ .  $I_{CEO}$  is collector-emitter leakage with the base open and defines the Common Emitter Breakdown Voltage,  $BV_{CEO}$ .  $I_{CBO}$

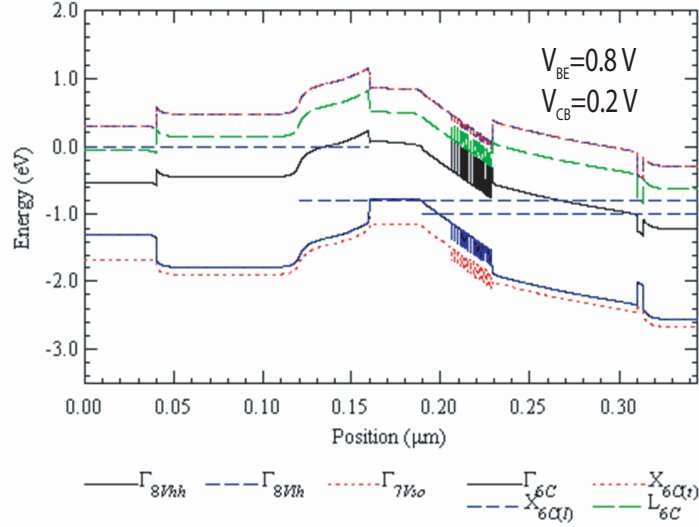
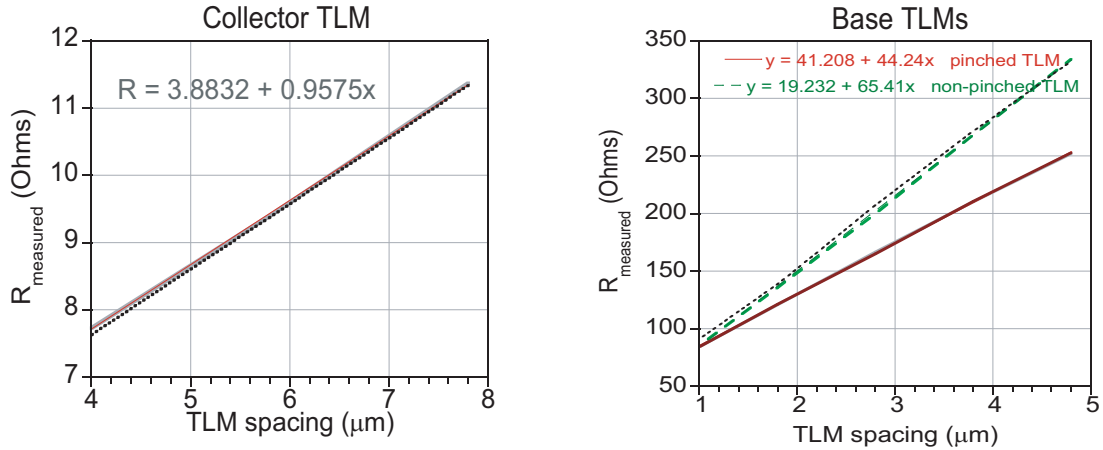


Figure 4.15: Band diagram of Implanted subcollector DHBT at  $V_{be} = 0.8\text{V}$  and  $V_{cb} = 0.2\text{V}$

is the collector-base diode leakage with the emitter open and defines the Common Base Breakdown Voltage  $BV_{CBO}$ . It can be shown that  $I_{CEO} = \beta I_{CBO}$  and hence  $BV_{CEO}$  is usually smaller than the common base breakdown  $BV_{CBO}$ .

Typically breakdown voltages are defined at a certain current, say  $I_{CBO} = 50 \mu\text{A}$ . By this definition,  $BV_{CEO}$  (at  $I_c = 50 \mu\text{A}$ ) = 4.3V and  $BV_{CBO} = 5.2\text{V}$ . As detailed in Chapter 2, with  $V_{be}$  spacings of 1mV and  $V_{cb} = 0\text{V}$  and 0.1V, the resistance as seen in the emitter  $R_{ex} + R_{bb}/\beta$  is derived from gummels and plotted as function of current density Fig. 4.21. The decrease at low current densities ( $\approx 1 - 3 \text{ mA}/\mu\text{m}^2$ ) is an artifact due to the exponential curve fitting to  $I_c$ . However at medium current densities,  $R_{ex} + R_{bb}/\beta$  is seen to decrease. This is due to two effects- the increase in current gain and more importantly decrease of base resistance at medium-high current densities. At these current densities, the base current is partly carried by electrons [20]. The thermal resistance can also be derived. Plotted in the right hand side of Fig. 4.21 is the thermal resistance,  $\Theta_{th} \cdot \phi$  (also referred to as  $R_{th}$  are derived. The thermo electric feedback coefficient  $\phi$ , for InP/InGaAs DHBT, at  $1 \text{ mA}/\mu\text{m}^2$  is  $\approx 0.001 \text{ mV}/^\circ\text{C}$  [21]. From this the thermal resistance at  $1 \text{ mA}/\mu\text{m}^2$  is 2.1 deg K/mW. The collector resistance is extracted from the  $I_C - V_{CE}$  characteristics of the transistor at saturation, using the Fixed  $\beta$  method described in Chapter 2 and is plotted in Fig. 4.22. This method returns the collector plus emitter series

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Layers	$R_{sh}$ ( $\Omega/sq$ )	$\rho_c$ ( $\Omega-\mu m^2$ )	$\rho_{horz}$ ( $\Omega-\mu m$ )	$L_T$ ( $\mu m$ )
Base	1100	50	115	0.21
Collector	24	100	100	2.04

Figure 4.16: Base and Collector TLM measurements

resistance. Since the emitter resistance is already known,  $R_{ex} \approx 3 \Omega$  the collector resistance is determined to be  $\approx 12 \Omega$ . The device DC current gain, ideality factors, and collector leakage current are consistent with those measured from the triple-mesa HBT equivalent [3].

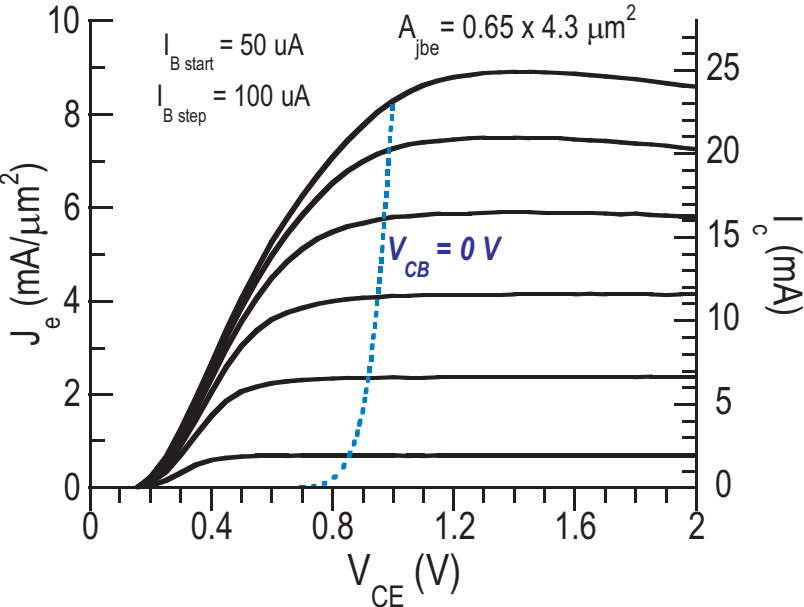


Figure 4.17: DC  $I_C - V_{CE}$  characteristics

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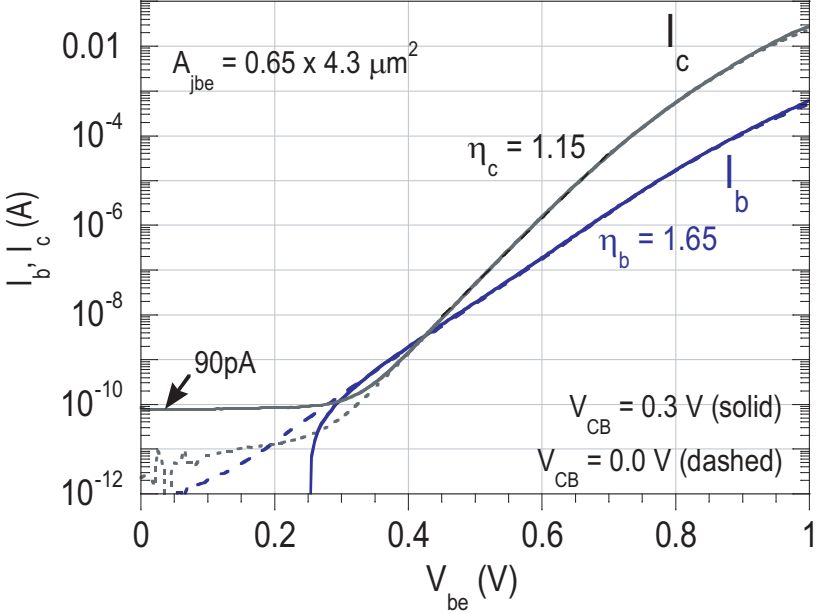


Figure 4.18: Gummel curves at  $V_{cb} = 0\text{V}$  and  $0.3\text{V}$

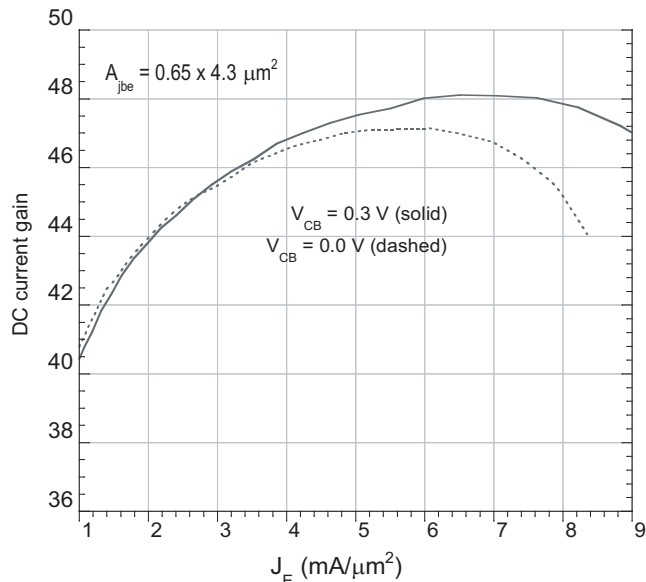


Figure 4.19: DC gain of the HBT at  $V_{cb} = 0\text{V}$  and  $0.3\text{V}$

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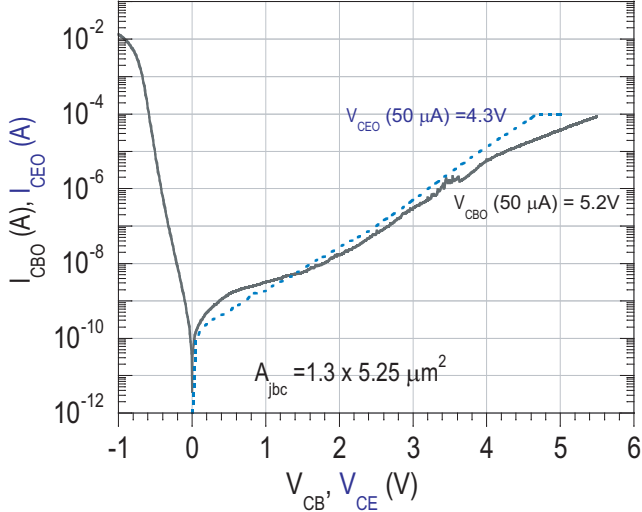


Figure 4.20: Leakage currents in the device

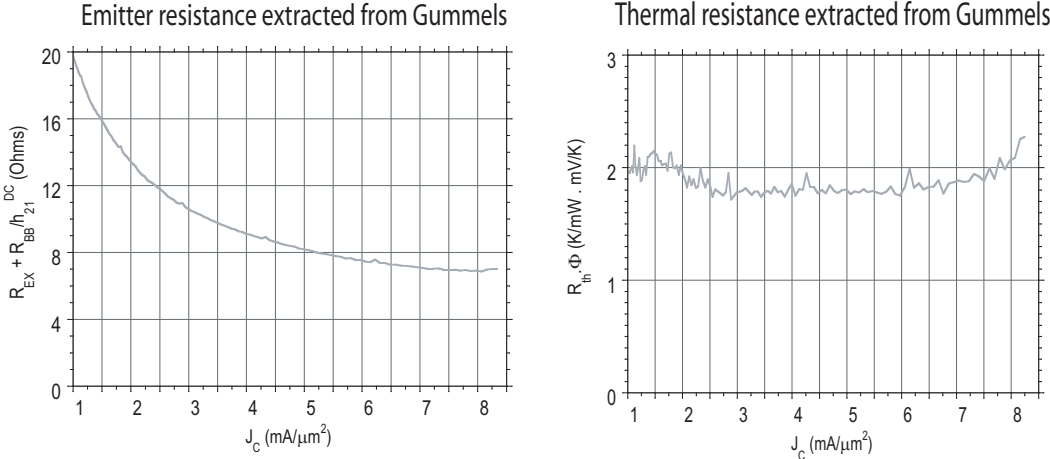


Figure 4.21: Extraction of Emitter resistance, and thermal resistance



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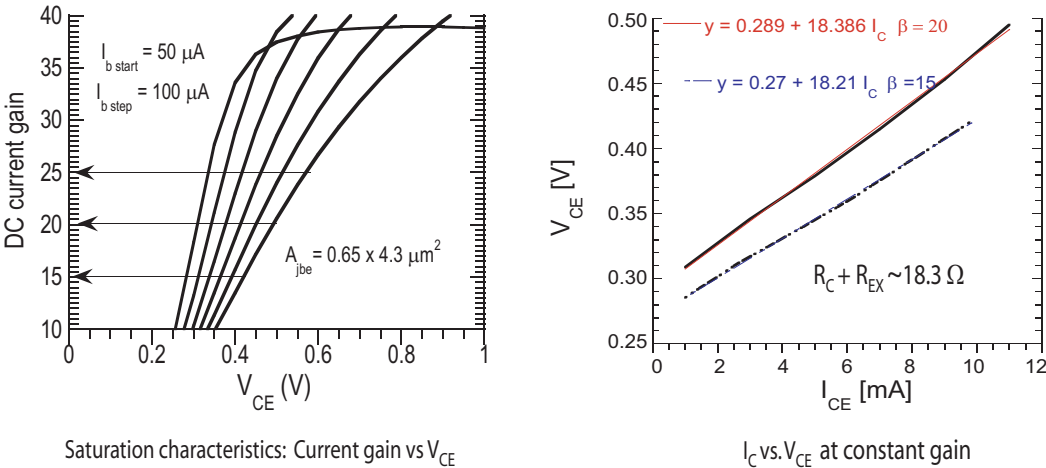


Figure 4.22: Extraction of Collector resistance

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DC-45 GHz S-parameter measurements are carried out after performing an off wafer Line-Reflect-Reflect-Match calibration on an Agilent 8510C network analyzer. On-wafer open and short circuit pad structures identical to the ones used by the devices are measured after calibration in order to de-embed their associated parasitics from the device measurements. Shielded *Infinity* probes are used for microwave measurements. Their superior field confinement reduces unwanted couplings to nearby devices and transmission modes. The unilateral power gain,  $U$  and short circuit common emitter current gain,  $h_{21}$  are plotted in Fig. 4.23. Extrapolating these at 20 dB/decade,  $f_{\tau}$  and  $f_{max}$  are obtained as the frequencies at which  $U$  and  $h_{21}$  are 0 dB respectively. The  $f_{max}$  and  $f_{\tau}$  are lower than the standard, fully

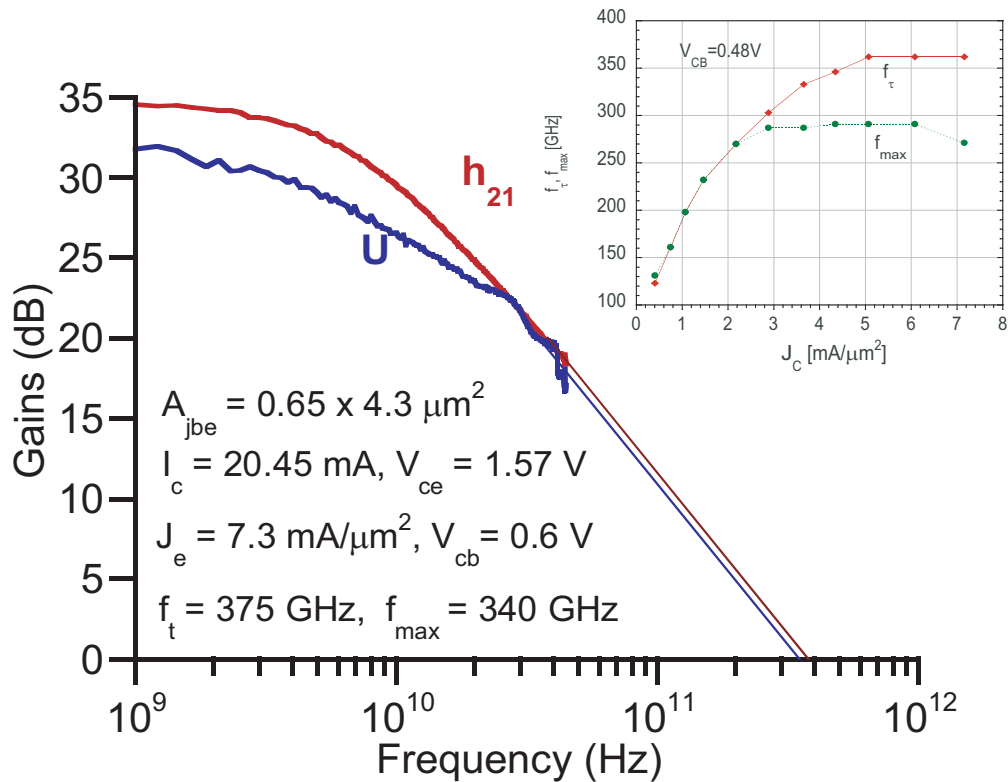


Figure 4.23: Microwave gains of Si implanted subcollector DHBTs

epitaxial, triple mesa DHBT. The base resistance is higher for these DHBTs ( $45 \Omega$  vs.  $35 \Omega$ ), since these were grown at UCSB where there are difficulties obtaining a high Carbon doping and hence lower base resistivity. Further, the collector resistance is higher ( $10 \Omega$  vs.  $2 \Omega$ ) due to the poor ohmic contact. The capacitance voltage

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characteristics of the implanted subcollector DHBTs at  $I_c = 0$  mA, are plotted and compared with the standard mesa equivalent in Fig. 4.24. From the figure, contrary to the expected reduction, the  $C_{cb}$  is the same as that of the standard triple mesa device lower voltages. The  $C_{cb}$  is the same for the standard mesa device and the implanted subcollector DHBTs till  $\sim 0.6$ V.  $C_{cb,pad}$  is thus not eliminated. As seen from the CV plots, the capacitance of the implanted subcollector DHBTs where the pad is isolated progressively decreases with increasing bias. This is not seen in the triple mesa equivalent where the collector is fully depleted at  $\sim 1$ V and the reduction in capacitance due to the Debye tail is negligible. In order to investigate this phenomenon, CV measurements ( $I_c = 0$  mA) are made of devices where the base access pad is not isolated and plotted in Fig. ???. The devices where the base access pad is isolated is similar to a standard mesa DHBT with a base access pad capacitance, since the Si implant is also performed in the regions under the base pad. It is seen from Fig. ??? that this capacitance decrease at high bias voltages is not observed for implanted subcollector DHBTs where the base pad is not isolated and closely follows the CV profile of the standard mesa DHBT. This data suggests the presence of some interface charge under the base access pad. This charge is progressively depleted at high collector base bias voltages and hence a reduction is seen in devices where the pad is isolated as in Fig. 4.1. The DC results compare with the standard mesa DHBT and hence indicate the feasibility of manufacturing microwave HBTs on implanted substrates. The RF results indicate that the expected  $C_{cb}$  reduction due to isolation of base access pad by implant is not present at low bias voltages. The capacitance due to the base access pad,  $C_{cb,pad}$  is thus not eliminated at these voltages. This is attributed to an N-type charge at the growth interface which is depleted at progressively higher biases. The elimination of this interface charge is therefore crucial to the theoretically predicted operation of these implanted subcollector DHBTs.

### 4.5 Interface charge

It was seen from the previous section that there exists a positive interface charge that acts as terminating electrode for the base-collector electric field. This charge, if left uncompensated, results in incomplete depletion and hence no reduction in extrinsic  $C_{cb}$ .  $C_{cb,pad}$  is thus still present in this bias range (till  $\sim 3$ V). The interface charge is depleted at high bias voltages ( $\sim 4$ V) and hence there is a reduction in capacitance at those voltages. However, this is not very useful for bandwidth enhancement in transistor operation in InP based circuits.

The origin of this unintentionally introduced N-type conduction in exposed InP surfaces(epitaxial or substrate) is identified to be Si impurity states [22, 23]. These

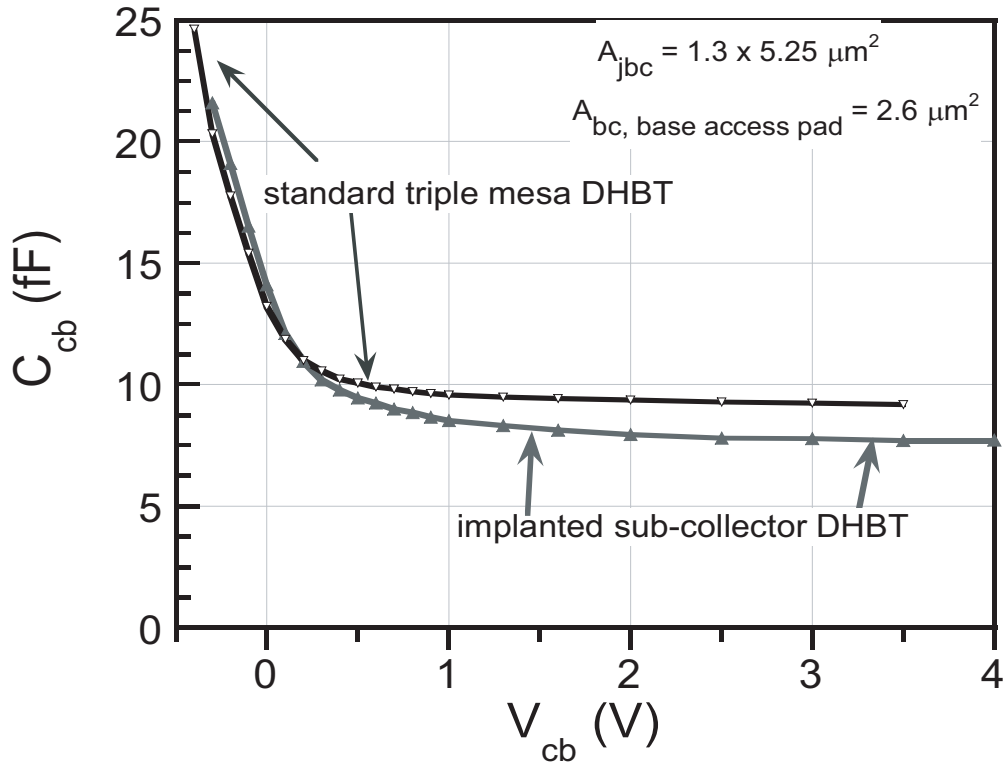


Figure 4.24: Capacitance voltage characteristics, at  $I_c = 0$  mA, of implanted sub-collector DHBTs with the base pad isolated

are verified by sheet charge measurements and SIMS. The Si donor states are formed as result of processing and also due to the ambient Si. The Si atom sheet concentration was found to be proportional to the time of exposure to ambient air. The concentration of these Si donor states is thus heavily process dependent and is in the range of  $1 \times 10^{12} - 1 \times 10^{13} \text{ cm}^{-2}$ . This interface charge is also responsible for parallel conduction at the substrate interface in InP based HFETs. It is the cause of the low turn on, increased resistance and high capacitance in p-i-n InP based photodetectors, where the p layer is regrown and the interface charge is present on the surface of the i-layer. Regrown HBTs with implanted pedestals reported at UCSB [24] also suffer from this charge at the regrowth interface which resulted in no capacitance reduction at low bias voltages.

Several methods have been utilized for reduction of this interface charge, including advanced surface clean procedures [23], annealing in a phosphine ambient [22]

and overgrowth of a charge compensation layer [25]. This interface charge is process dependent and full compensation might not be possible with process techniques or an empirical  $p^{++}$  compensation layer.

#### 4.5.1 Insufficient charge compensation with a $p^{++}$ overgrowth

The interface charge is also present on epitaxially grown UID InP layers, as part of the pedestal process developed at UCSB [25]. Starting with a grown template of 200 nm UID InP over 300 nm  $N^{++}$  InP, Si is selectively implanted to form an  $N^{++}$  pedestal. The active collector, base and emitter layers are then regrown. Interface charge is present on the 200 nm UID epitaxial InP prior to regrowth and has the same effect as here. An empirically determined  $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  compensation layer was overgrown. In order to check the level of  $p^{++}$  doping required for compensation, various levels of  $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  from  $2.5\text{-}7.5 \times 10^{18} \text{ cm}^{-3}$ , and a test sample with undoped  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$ , are first overgrown on the template. This is followed by growth of the active device layer stack - the 120 nm  $N^-$  drift collector, base and emitter. This device has a 120 nm collector on UID InP and with sufficient compensation has a depletion depth of 320 nm. This is identical to the extrinsic region of the device in the regrown pedestal process [25]. Large area HBTs are fabricated and Capacitance - Voltage measurements indicate the extent of interface charge compensation. Fig. 4.25 shows the CV plots of the extrinsic device region with various levels of  $p^{++}$  compensation layers. As a result of interface charge, the layer without a  $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  layer has the largest capacitance at low bias voltages. The  $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  layer doped, in different samples, from  $2.5\text{-}7.5 \times 10^{18} \text{ cm}^{-3}$  incompletely compensate the interface charge as seen in the partial reduction in capacitance. The capacitance at 1V, for the layers without any  $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  overgrowth is  $\sim 33$  fF while the The capacitance at 1V, for the layers with 4 nm,  $7.5 \times 10^{18} \text{ cm}^{-3}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  overgrowth is  $\sim 23$  fF. Since the collector thickness is 120 nm and the thickness of the UID InP layer is 200 nm, if the interface charge is fully compensated, the expected capacitance when the collector is fully depleted is  $\sim 13$  fF. The tail in the CV at high biases indicates that the interface charge is insufficiently compensated and gets progressively depleted with increasing base collector voltage. As seen from the charge profile extracted from the CV data, there is a charge buildup at the growth interface, despite the 4 nm,  $5 \times 10^{18} \text{ cm}^{-3}$   $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  compensation layer. These results indicate that the empirical  $p^{++}$  layer is not a good solution to compensate the interface charge.

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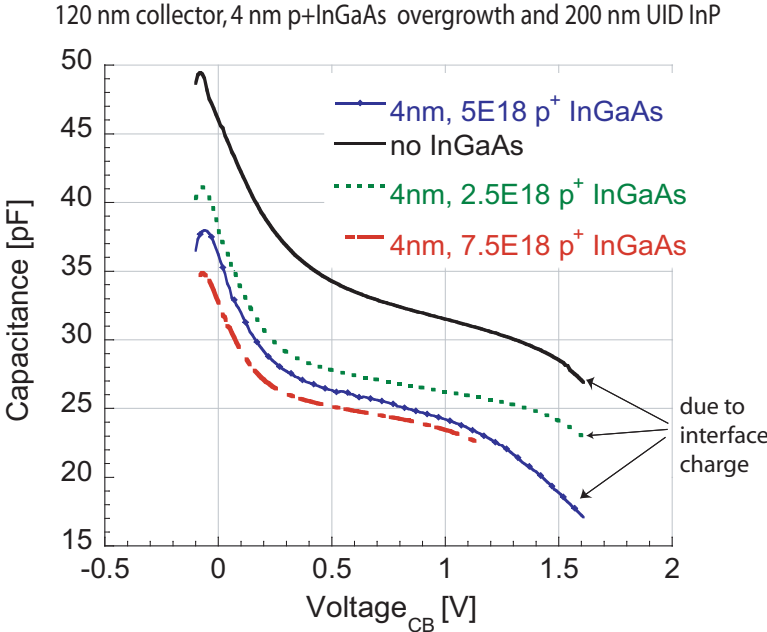


Figure 4.25: C-V measurements of HBTs where active device layers are grown over a 200 nm UID InP/300 nm  $N^{++}$  InP template. Various levels of  $p^{++}$   $In_{0.53}Ga_{0.47}As$  layers are overgrown as the first layer on the template to determine the  $p^{++}$  doping required to compensate interface charge

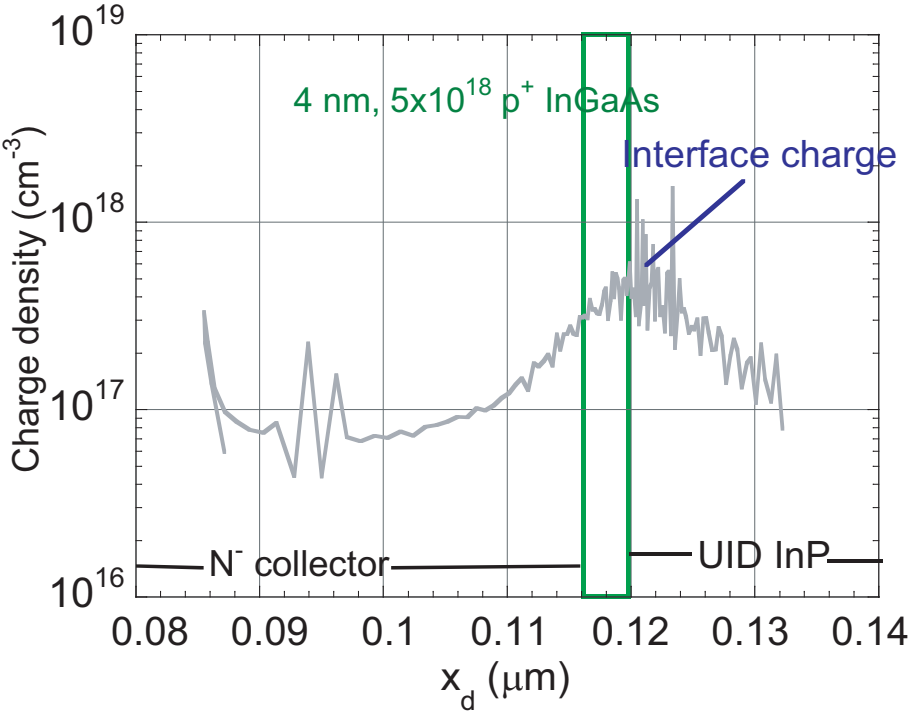


Figure 4.26: Interface charge density on 200 nm UID InP/300 nm  $N^{++}$  InP template with 4 nm of  $5 \times 10^{18} \text{ cm}^{-3} p^{++} \text{ In}_{0.53}\text{Ga}_{0.47}\text{As}$  overgrowth

### 4.5.2 Interface charge compensation using Fe implant

A new, robust technique is introduced in this dissertation wherein Fe implantation is used to eliminate this interface charge. As discussed in Chapter ?? , Fe is a mid-gap acceptor in InP and serves to compensate the donor states. Thus by sufficiently compensating the Si donor states with Fe, the N-type conduction at the interface is eliminated. The process thus involves, implanting a blanket sheet of Fe to compensate the interface charge. The subsequent selective Si subcollector implant then defines the isolated subcollector as in §??

The band diagram of the  $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$  and  $N^-$  collector region under the base pad is shown in Fig. ?? when there is an N-type interface present and the corresponding simulated CV profiles are shown in Fig. ?. To first order, the electric field in the collector corresponds to,

$$\frac{\epsilon(V_{cb} + \phi_{bi})}{qT_c} = n_s + \frac{N_c T_c}{2} \quad (4.5.0)$$

where  $n_s$  is the regrowth interface charge density,  $N_c$  is the doping in the drift collector, and  $T_c$  is the thickness of the drift collector,  $\phi_{bi}$  is the built-in potential of the base-collector junction and  $V_{cb}$  is the base-collector reverse bias voltage. In order to ensure complete charge depletion at the regrowth interface in the region under the base pad area

$$\frac{\epsilon(V_{cb} + \phi_{bi})}{qT_c} + n_{Fe} \geq n_s + \frac{N_c T_c}{2} \quad (4.5.0)$$

where  $n_{Fe}$  is the density of active Fe in  $\text{cm}^{-2}$ . If the collector is so designed as to be fully depleted at zero bias, then

$$\frac{\epsilon\phi_{bi}}{qT_c} = \frac{N_c T_c}{2} \quad (4.5.0)$$

Therefore  $n_{Fe} \geq n_s$  for full charge compensation, even at zero bias. The Fe implant conditions are chosen to satisfy the above requirement. The conditions for the Fe implant are as follows

- The Fe implant conditions should be selected to fully compensate the N-type charge at the growth interface between the substrate and collector epitaxial layers, over the observed  $1 \text{ } 5 \times 10^{12} \text{ cm}^{-2}$  range of regrowth interface charge densities.
- Some of the surface is unintentionally attacked during various process steps. The Fe implant should be therefore be sufficiently deep.

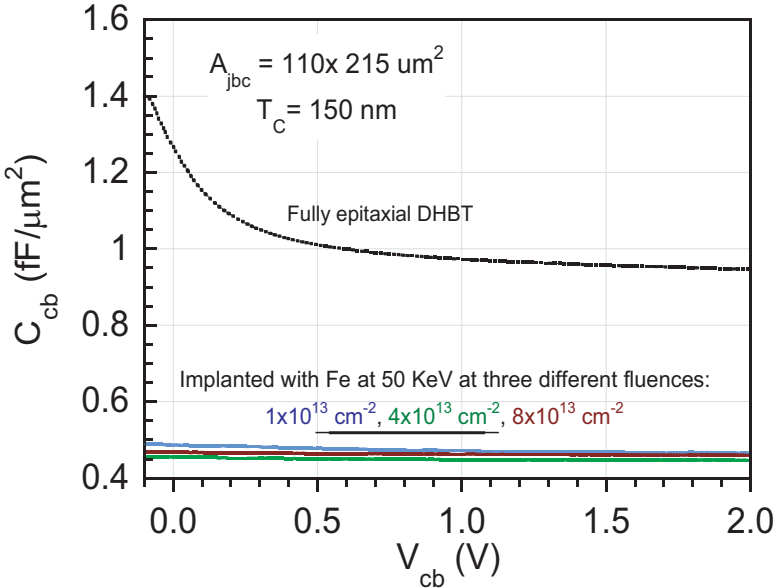


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- Fe is a heavy ion and causes significant damage to the crystal. This should be completely annealed out before growth.
- The Fe implant should not be very high since the subsequent Si implant should overcome the Fe doping to form a highly  $N^{++}$  subcollector for the active device regions.

A experiment, similar to the one in §4.5.1, to check C-V curves of large area devices is performed to verify the compensation of interface charge by Fe. In order to determine the Fe doping required for complete compensation, Fe is implanted, at 50 KeV at three different fluences in a similar grown template, 200 nm UID InP over 300 nm  $N^{++}$  InP. These were then annealed at 700 °C for 5 minutes, which is shown in Chapter 5 to be sufficient to repair the lattice damage due to Fe implant for fluences below  $2.4 \times 10^{14}$  ions/cm<sup>2</sup>. The active layers (with a 150 nm collector) are immediately grown after the Fe implant. Since the Fe implant is intended to compensate the interface charge, these do not have a compensating  $p^{++}$  overgrowth. Large area devices are fabricated to determine C-V characteristics. A fully epitaxial, standard DHBT with similar active layer structure is also co-processed for comparison. From Fig. 4.27 it is seen that the capacitance is reduced over the entire measured range of bias voltages, for all fluences of Fe. Thus the interface charge is fully compensated even for a fluence as low as  $1 \times 10^{13}$  cm<sup>-2</sup>. Capacitance when collector is fully depleted for fully epitaxial DHBT is  $1 \text{ fF}/\mu\text{m}^2$ . Capacitance when collector is fully depleted for Fe implanted, regrown DHBT is  $0.42 \text{ fF}/\mu\text{m}^2$ . This corresponds to a depletion depth of  $\approx 350$  nm.  $T_c(150\text{nm}) + T_{undopedInP}(200\text{nm}) = 350$  nm. The depletion depth is increased and capacitance is therefore reduced, exactly as expected in the extrinsic device regions.

Fig. 4.28 and Fig. 4.29 shows further evidence of compensation of interface charge by Fe implant at two different fluences. In LACVD 3 and LACVD 6, Fe is selectively implanted. Since Fe is present only in the extrinsic region, current blocking effects (if any) due to Fe traps are removed. In both figures, LACVD1 and LACVD2 have similar profiles which is again proof that Si doping is sufficiently high to counter compensate Fe that no shift in built in voltages are observed. In LACVD5 and LACVD6, the effect of interface charge is observed, where the capacitance starts to decrease at  $\sim 2\text{V}$  of reverse bias.



Layers	Energy	Dose
150 nm collector	50 KeV	$1 \times 10^{13} \text{ ions/cm}^2$
Fe implant		$4 \times 10^{13} \text{ ions/cm}^2$
UID InP 200nm		$8 \times 10^{13} \text{ ions/cm}^2$
N++ InP 300nm		

All implants are at 200°C at an offset of 7°  
 Anneal is 700 °C for 5 minutes

Figure 4.27: CV measurements of pedestal templates implanted with varying degrees of Fe compared with a standard, fully epitaxial DHBT

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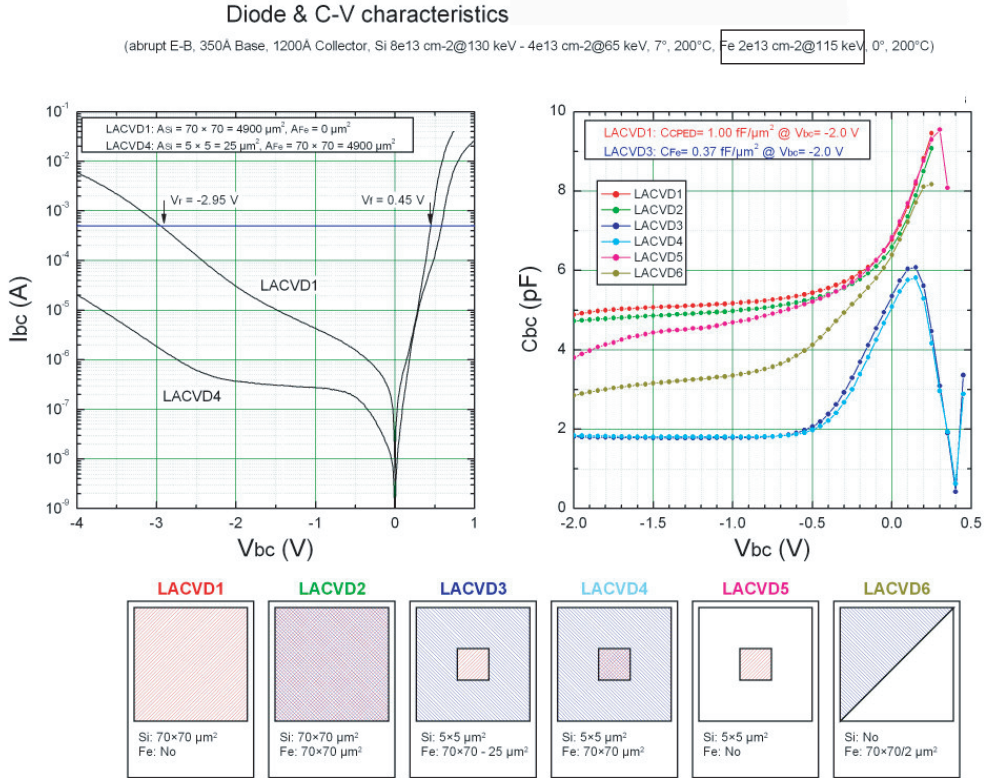


Figure 4.28: C-V's of various test structures: Fe is implanted at  $2 \times 10^{13}$  ions/cm<sup>2</sup>: This data was obtained at RSC

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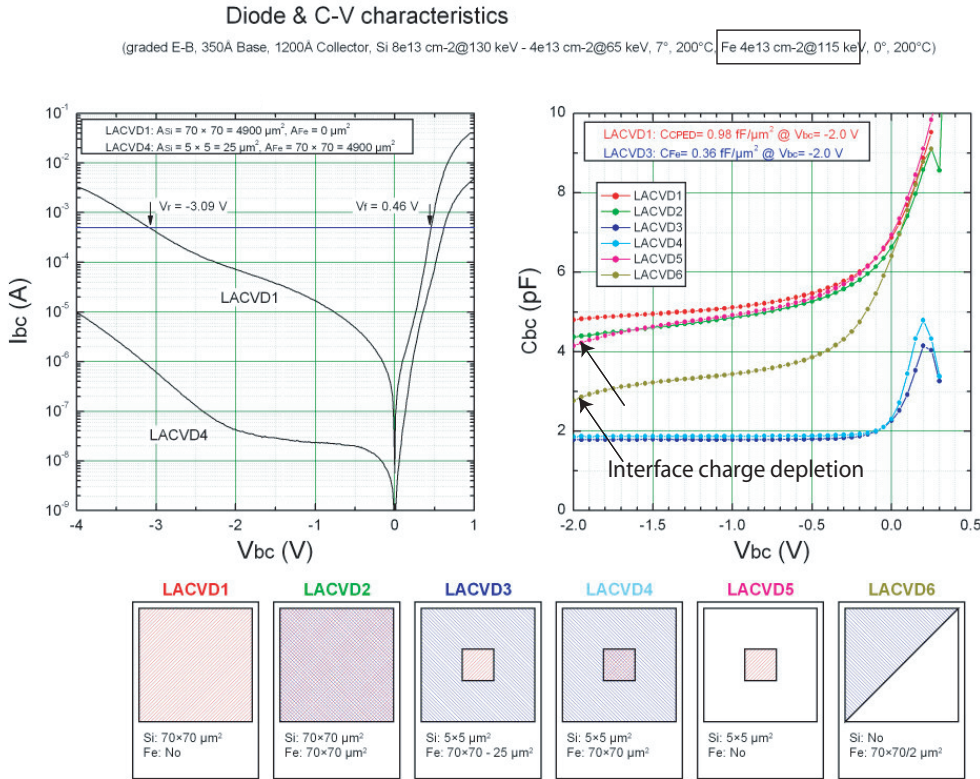


Figure 4.29: C-V's of various test structures: Fe is implanted at  $4 \times 10^{13}$  ions/cm $^2$ : This data was obtained at RSC

### 4.5.3 Large area devices with Si and Fe implants

The selective Si subcollector implant that follows the blanket Fe implant must dope the region  $N^{++}$ . Large area HBTs are fabricated on templates with and without Si implant. On one set of templates, the active layers (with a 150 nm collector) are immediately grown after a 50 KeV,  $2 \times 10^{13}$  ions/cm<sup>2</sup> Fe implant. It was seen from the previous experiment that any fluence  $\geq$  is sufficient for interface charge compensation. On another set of templates, following the Fe implant, a blanket Si implant similar to the one used for the implanted subcollector DHBT, is carried out and the active device stack is then grown. A fully epitaxial, standard DHBT with similar active layer structures is also co-processed as a comparison. Fig. 4.30 shows the C-V profile of the three different test structures. The std. mesa DHBT with 150 nm collector has a capacitance of  $\approx 0.8$  fF/ $\mu\text{m}^2$  when fully depleted. The HBT overgrown on Fe implanted template has a capacitance of  $\approx 0.34$  fF/ $\mu\text{m}^2$ . In the intrinsic region under the emitter, the  $N^+$  pedestal is present. This is exactly represented by the large area HBT where both Si and Fe are implanted prior to growth. The CV characteristics where Si and Fe are implanted, in Fig. 4.30, is similar to the standard mesa DHBT. This is to be expected, since a blanket  $N^+$  pedestal layer is akin to a  $N^{++}$  subcollector layer. The built-in voltage of the base collector junction of the pedestal HBT is similar to the fully epitaxial HBT. This is the expected performance in the intrinsic portion of the device.

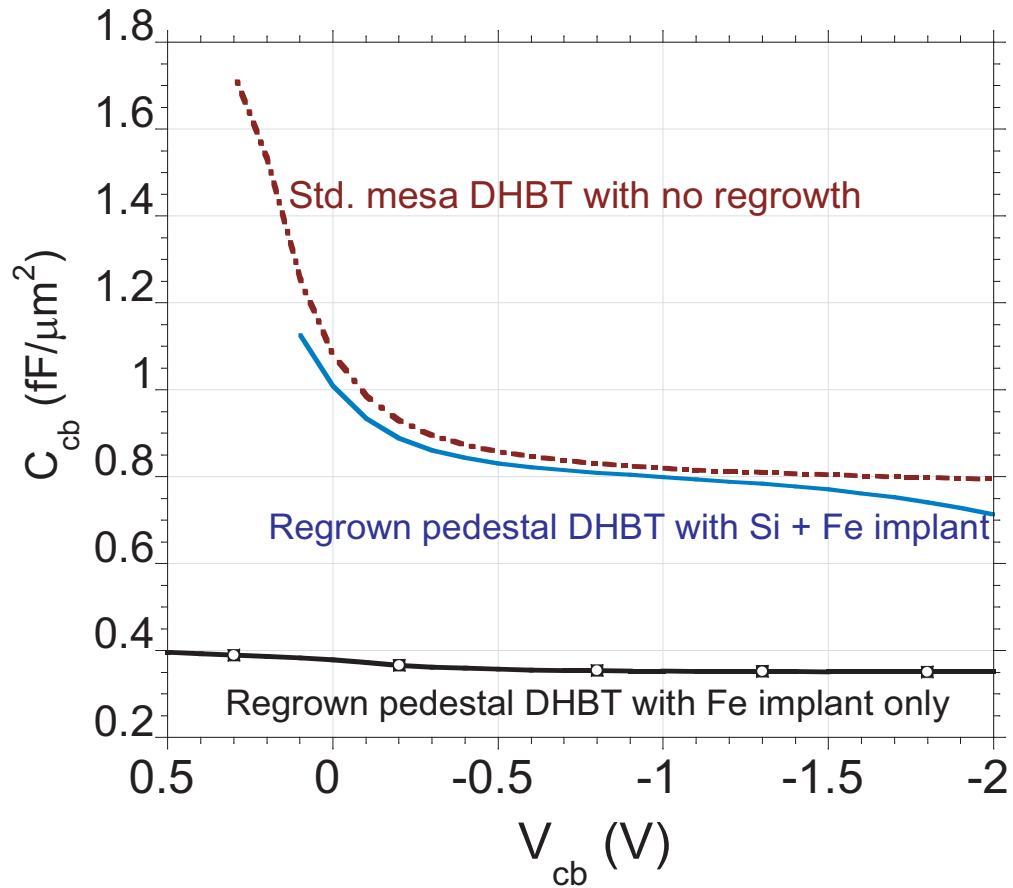


Figure 4.30: C-V characteristics of standard fully epitaxial HBT, regrown HBT with only Fe implant on 200 nm UID InP/ 300 nm  $N^{++}$  InP template, and regrown HBT with Fe and Si pedestal implants on 200 nm UID InP/ 300 nm  $N^{++}$  InP template

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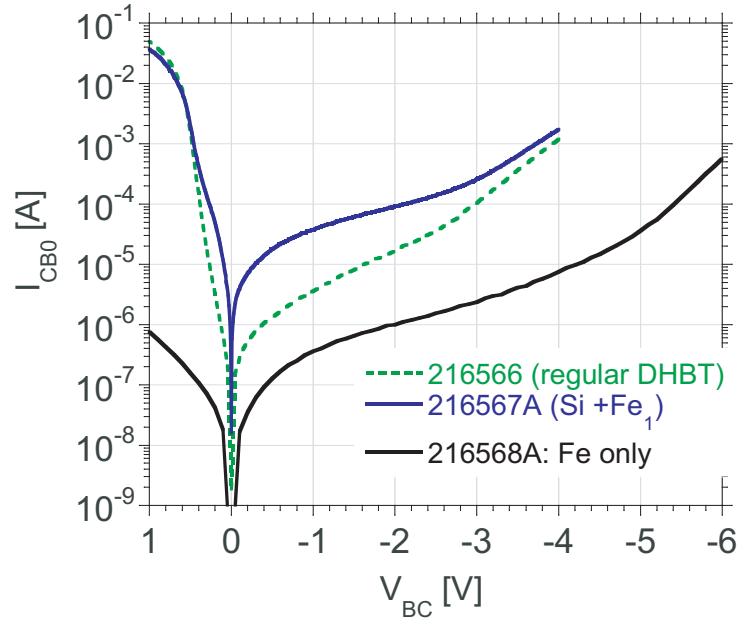


Figure 4.31: Comparison of reverse base collector leakage characteristics of std. mesa HBT, regrown HBT with Fe implanted on 200 nm UID InP/ 300 nm  $N^{++}$  InP template, and regrown HBT with Fe and Si pedestal implants on 200 nm UID InP/ 300 nm  $N^{++}$  InP template

The base collector leakage for the regrown device with Si and Fe implants is similar to a fully epitaxial HBT. The base collector leakage plot shows that  $I_{CBO}$  at 3V  $\sim 0.1$  mA ( $10 \times 10^{-6}$  mA/ $\mu\text{m}^2$ ). For a microwave device with a base collector junction of  $\sim 7 \mu\text{m}^2$ ,  $I_{CBO}$  and gain of  $\sim 50$ ,  $I_{CEO}$  is  $\approx 3.5 \times 10^{-3}$  mA/ $\mu\text{m}^2$ . This is a very small leakage current at 3V. The DC  $I_C - V_{CE}$  characteristics do not show any current blocking due to the Fe implant. The offset voltages and breakdown voltages are consistent with those measured for similar large area, fully epitaxial HBTs. There is a high series collector resistance which is possibly due to low doping in the pedestal (Si is implanted at 150 KeV /  $2 \times 10^{13}$  ions/cm $^2$  to minimize implant straggle which will be addressed in Chapter ?? ).

Low collector contact resistivity and sheet resistance are obtained in §?? even with an Fe implant. Therefore the Si implant sufficiently overcomes the Fe implant to form an  $N^{++}$  subcollector (or pedestal) and does not hamper the collector ohmic contact.

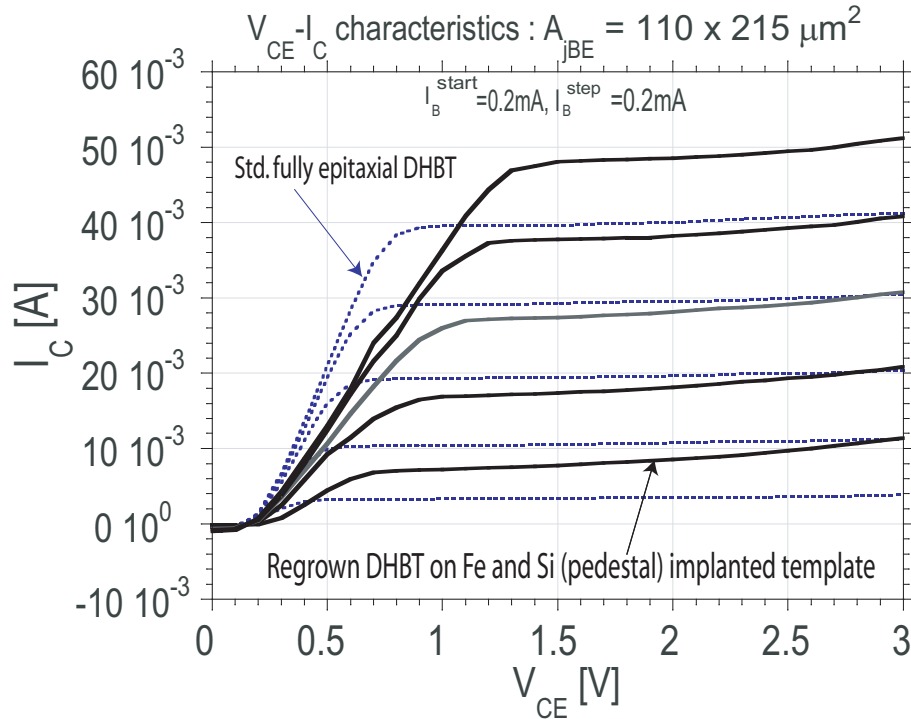


Figure 4.32: DC characteristics of regrown HBT with Fe charge compensation implant and Si pedestal implants on 200 nm UID InP/ 300 nm  $N^{++}$  InP template. The device characteristics of the standard fully epitaxial DHBT is shown for comparison

## 4.6 Implanted subcollector DHBTs with Fe

The process is modified to have include a blanket shallow Fe implant. It was seen from the previous section, that Fe implant with fluences  $\geq 1 \times 10^{13}$  ions/cm<sup>2</sup> is sufficient to compensate the residual interface charge. Fig. ?? shows the profile of the Fe at the implant conditions chosen, and also the Si subcollector implant. A low energy Fe is chosen to minimize current blocking effects and to ease the requirements on Si implants. Assuming  $\sim 10$  nm of the surface is removed, the active Fe concentration is  $\sim 1 \times 10^{13}$  cm<sup>-2</sup>. This is much larger than the observed range of interface charge densities. The Si doping at the surface is designed to be high enough to enable a low collector contact resistance.

The process flow of the implanted subcollector HBT with Fe is shown in Fig. 4.33. A semi-insulating InP substrate is blanket implanted with Fe at 10 keV, at a fluence of  $2 \times 10^{13}$  ions/cm<sup>2</sup> as in Fig. ??a. The wafer is then annealed in the RTA at 700



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°C for 5 minutes to activate the Fe and to repair crystal damage. The process flow that follows is identical to that of the implanted subcollector HBT without Fe, described in §???. TiW alignment targets are defined. The sample is then selectively implanted, using  $Si_xN_y$  as the implant mask, with Si. The Si and Fe implant conditions are given in Table 4.6. The Si implants are activated by annealing at 800°C/10s. The Si implant dose is much larger than the Fe acceptor density in the selectively implanted region. This ensures that a highly doped, isolated subcollector region is formed. The base access pad lies outside the subcollector implant boundary and, in the absence of interface charge, does not contribute to  $C_{cb}$ . The active InP HBT layers are then grown by MBE. The device drift collector, base, and emitter layers for both wafers are identical to that of the DHBTS reported in §4.4. As before, Growth is initiated with a 3.5nm undoped InGaAs etch-stop layer between the InP collector and the InP substrate. The drift collector, base, and emitter layers are grown. Devices are formed by wet-etching the emitter and base mesas. The collector contact used is Ni/Ge/Au/Ni/Au as in §4.1.2 and are annealed at 320 °C.

Table 4.6: Implant conditions for implanted subcollector HBT process with Fe

Implant species	Implant energy	Implant fluence	Offset angle	Implant temperature
Fe	10 KeV	$2 \times 10^{13}$ ions/cm <sup>2</sup>	7 °	200 ° C
Si	200 KeV	$3 \times 10^{14}$ ions/cm <sup>2</sup>	7 °	200 ° C
	40 KeV	$8 \times 10^{13}$ ions/cm <sup>2</sup>		
	10 KeV	$3 \times 10^{13}$ ions/cm <sup>2</sup>		
$I_{ions} \leq 10 \mu A$				

The sheet resistance of the subcollector with Si and Fe implant is determined from Hall measurements and shown in Table ??. The different anneal conditions

Table 4.7: Hall measurements of resistivity and mobility for Si implants

Anneal temperature (°C)	Anneal time (s)	Hall mobility (cm <sup>2</sup> /V · s)	Carrier density (cm <sup>-2</sup> )	Sheet resistance (Ω/□)
700	60	925	$2.6 \times 10^{14}$	26
750	60	956	$2.5 \times 10^{14}$	26
700	300	905	$2.5 \times 10^{14}$	27
750	300	872	$2.7 \times 10^{14}$	26
800	10	710	$3.3 \times 10^{14}$	27
800	30	886	$2.8 \times 10^{14}$	25

shown here, all yield a subcollector sheet resistance  $\sim 25$ -27 Ω/□. 800 °C for

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30 seconds was chosen as the activation anneal for all Si implants. Ignoring the effect of Fe, the activation is  $\approx 56\%$ . These are similar to the values obtained for the implanted subcollector DHBTS without Fe. The surface Fe does not affect the resistivity of the subcollector.

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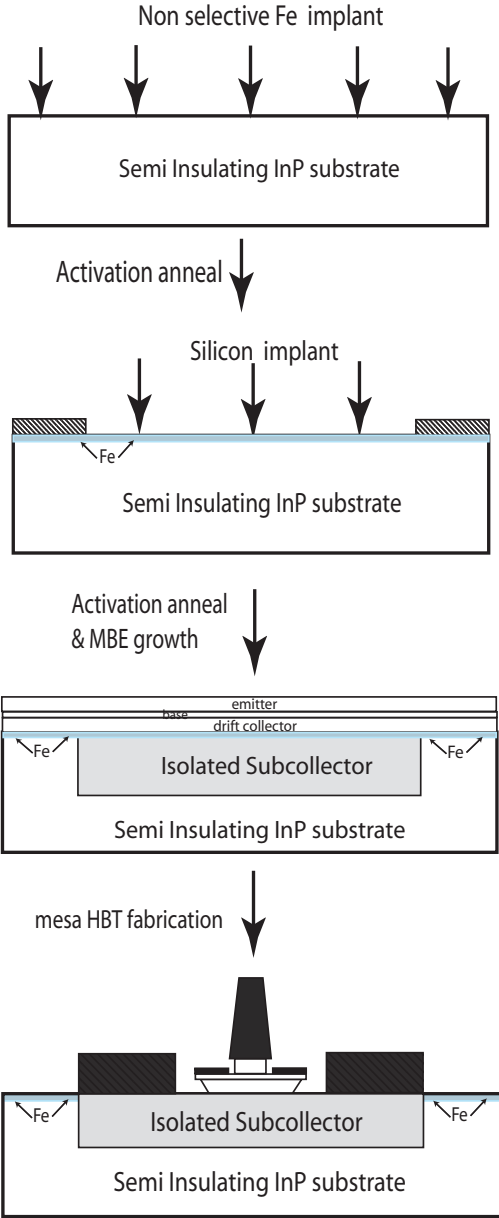


Figure 4.33: Process flow of the Implanted subcollector DHBT with Fe

## 4.7 Device Results - Si Implanted subcollector DHBTs with Fe

Large area devices have shown low leakage and excellent DC characteristics. Small area devices were processed with the identical layer structure shown in Table 4.5. The process has already been described in §4.4.1.

The TLM measurements of the collector and, of the pinched and non-pinched base TLMs are given in Fig. 4.34. The collector contact resistance is very high  $\approx 500 \Omega \cdot \mu\text{m}^2$ , while the sheet resistance is similar to that obtained in §4.4. Due to the  $320^\circ\text{C}$  anneal for the collector ohmics, the base ohmics also have high contact resistivity of  $\approx 165 \Omega \cdot \mu\text{m}^2$ . The Isolation between ohmic pads separated by  $10 \mu\text{m}$  is  $20 \text{ pA}/\mu\text{m}$ . Fig. 4.35 is a plot of the DC  $I_C - V_{CE}$  characteristics. There is no

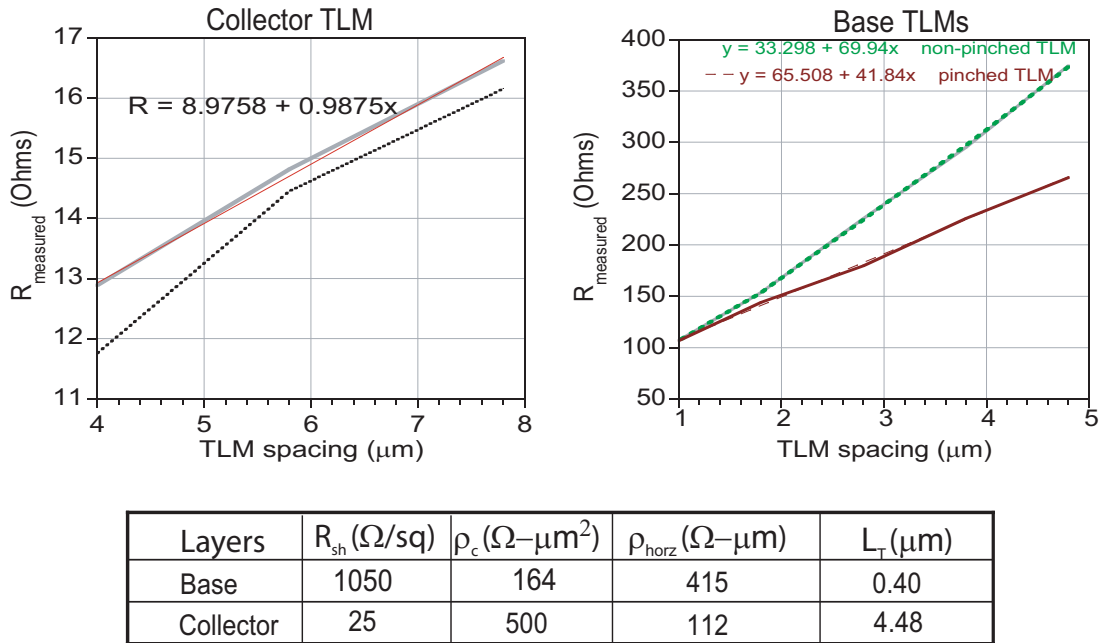


Figure 4.34: Base and Collector TLM measurements

evidence of gain compression till  $\approx 7 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0\text{V}$ . The Gummel curves are plotted for  $V_{cb} = 0\text{V}$  and  $V_{cb} = 0.3\text{V}$  Fig. 4.36. The collector and base ideality factors are  $\eta_c = 1.11$  and  $\eta_b = 1.58$ , consistent with those obtained for fully epitaxial DHBTs. The leakage current  $I_{CEO}$  from the gummel curves is  $\approx 1 \text{ nA}$  at  $V_{cb} = 0.3\text{V}$ . The DC current gain of the transistor  $= I_C/I_b$  from the gummel curves is shown in

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Fig. 4.37. The HBT's gain compression point is  $\approx 4 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0\text{V}$  and  $6 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0.3\text{V}$ . At high current densities,  $V_{CB,i} = V_{CB, applied} - I_C R_c$ . From the TLMs and the geometry of the transistor,  $R_c$  for this transistor is  $\approx 22 \Omega$ . At  $I_c \approx 15\text{mA}$ ,  $V_{CB,i} = V_{CB, applied} - 0.3\text{V}$ . The base collector junction is forward biased till  $V_{CB, applied}$  is  $\approx 0.3\text{V}$ . Shown in Fig. 4.38 are  $I_{CEO}$  and  $I_{CBO}$ . The

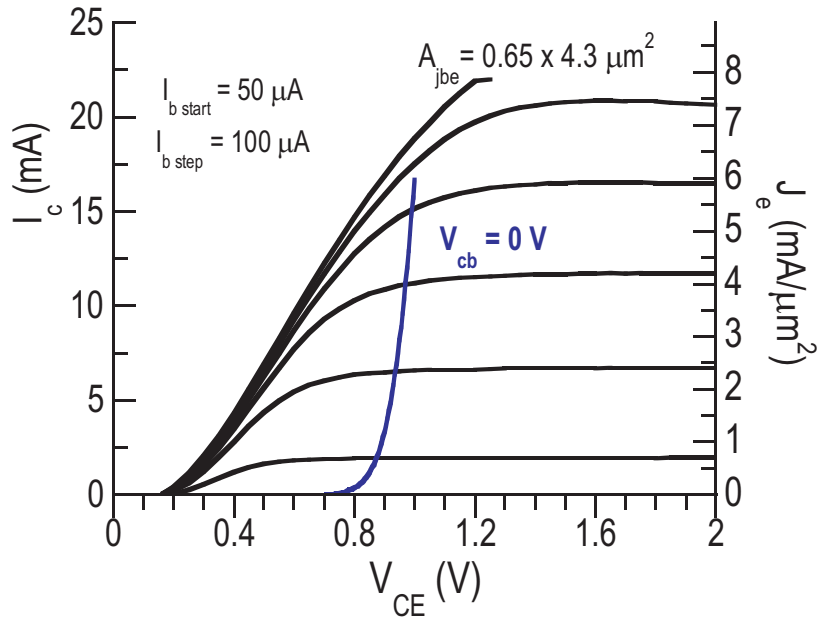


Figure 4.35: DC  $I_C - V_{CE}$  characteristics

breakdown voltages are defined at  $I_c = 1 \text{ mA}$ .  $BV_{CEO} = 5.6\text{V}$  and  $BV_{CBO} = 6.9\text{V}$ . For a fully epitaxial, standard mesa DHBT, the corresponding voltages are  $BV_{CEO} = 5.5\text{V}$  and  $BV_{CBO} = 6.8\text{V}$  at  $I_c = 1\text{mA}$ . The resistance as seen in the emitter,  $R_{ex} + R_{bb}/\beta$  is derived from gummels and plotted as function of current density Fig. 4.39.  $R_{bb}/\beta \approx 1 \Omega$  and therefore  $R_{ex} \approx 8 \Omega$ . Also plotted in Fig. 4.39 is the thermal resistance,  $\Theta_{th} \cdot \phi$  (also referred to as  $R_{th}$ ). The thermal resistance at  $5 \text{ mA}/\mu\text{m}^2$  is  $2.6 \text{ deg K}/\text{mW}$ . The collector resistance is extracted from the  $I_C - V_{CE}$  characteristics of the transistor at saturation, using the Fixed  $\beta$  method and is plotted in Fig. 4.40. The emitter resistance,  $R_{ex} \approx 8 \Omega$ , the collector resistance is determined to be  $\approx 22 \Omega$ , consistent with the value calculated from TLM measurements. This is very high compared to the value of  $1\Omega$  that is typically obtained and is due to the poor ohmic contact to  $N^{++}$  InP. Contact resistivities as low as  $7 \Omega \cdot \mu\text{m}^2$  have been obtained with ohmics to  $N^{++}$  InP. Therefore the ohmic contact process can be improved.

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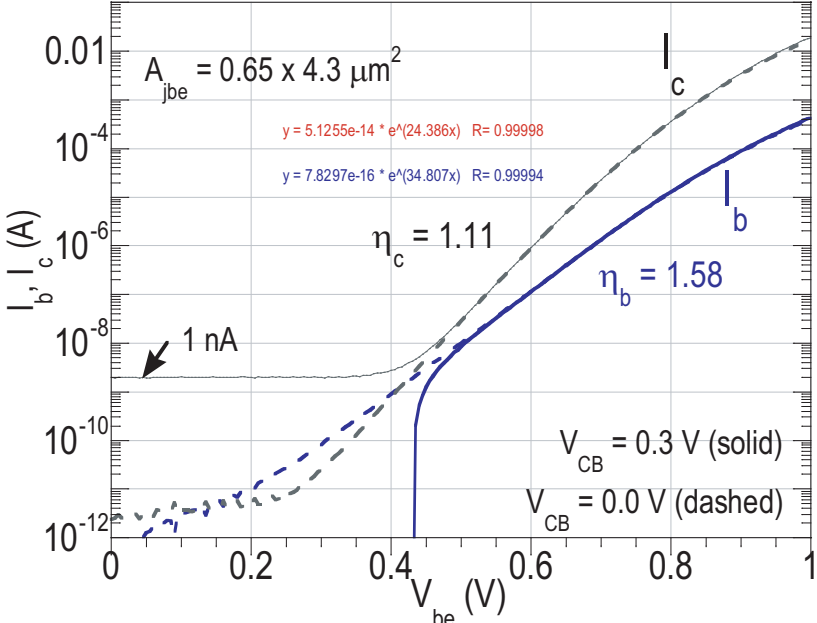


Figure 4.36: Gummel curves at  $V_{cb} = 0\text{V}$  and  $0.3\text{V}$

The device DC current gain, ideality factors, and breakdown voltages are consistent with those measured from the triple-mesa HBT equivalent [3].

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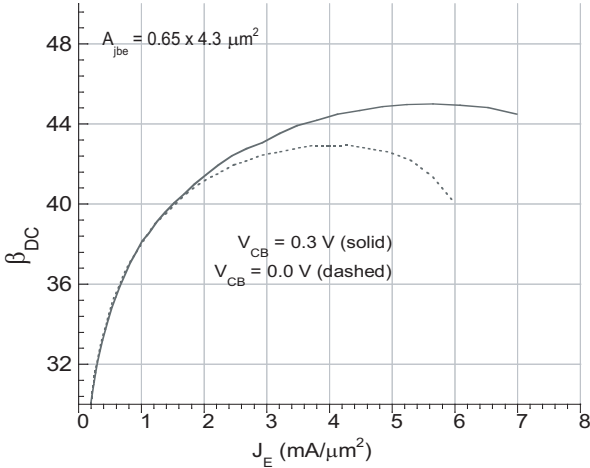


Figure 4.37: DC gain of the HBT at  $V_{cb} = 0\text{V}$  and  $0.3\text{V}$

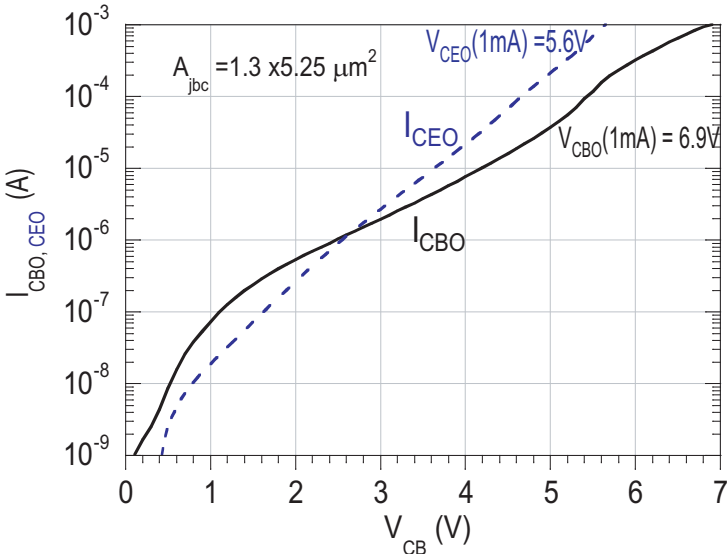


Figure 4.38: Leakage currents in the device

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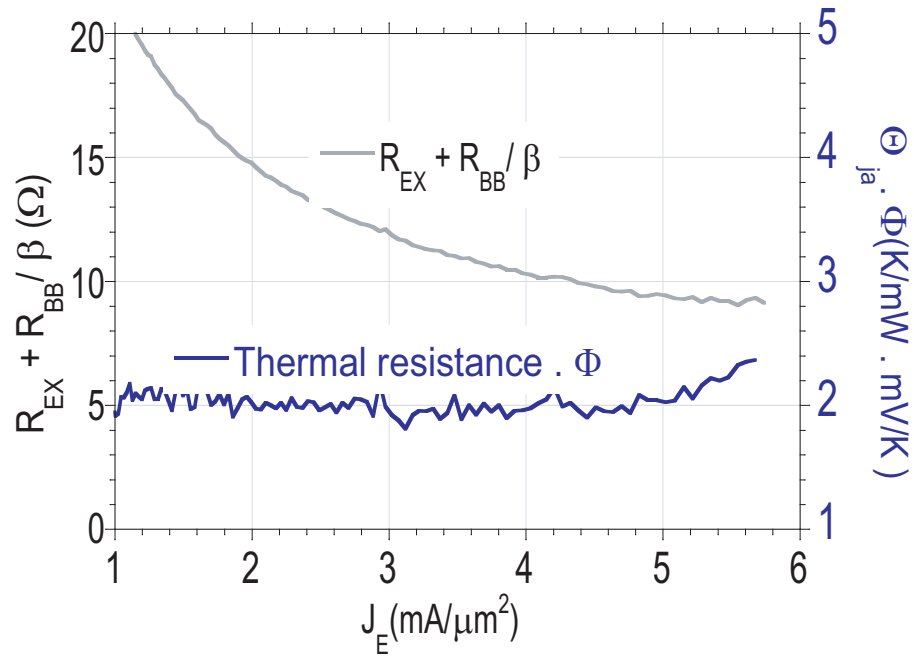


Figure 4.39: Extraction of Emitter resistance, and thermal resistance

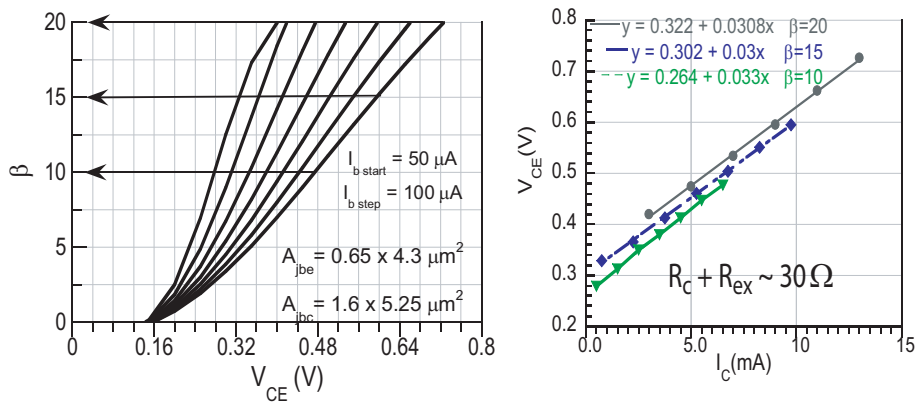


Figure 4.40: Extraction of Collector resistance



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DC-45 GHz S-parameter measurements are carried out after performing an off wafer Line-Reflect-Reflect-Match calibration on an Agilent 8510C network analyzer. On-wafer open and short circuit pad structures identical to the ones used by the devices are measured after calibration in order to de-embed their associated parasitics from the device measurements.  $f_{max}$  is determined from extrapolation through a least-square-fit between the transfer function,

$$U(f) = \frac{U_{DC}}{1 + U_{DC} \cdot (f/f_{max})^2} \quad (4.7.0)$$

to the measured microwave gain  $U$  at measured frequencies.  $1/f_\tau$  is the slope of  $\text{Im}(1/h_{21})$  [26] and is shown in Fig. 4.41. This method is more exact than a least square fit to,

$$|h_{21}(f)|^2 = \frac{1}{1/\beta^2 + (f/f_\tau)^2} \quad (4.7.0)$$

From these,  $f_{max} \approx 410$  GHz and  $f_\tau \approx 362$  GHz. These are lower than the standard, fully epitaxial, triple mesa DHBT. The base resistance is higher for these DHBTs (45  $\Omega$  vs. 35  $\Omega$ ), since these were grown at UCSB. Further, the collector resistance is much higher (22  $\Omega$  vs. 2 $\Omega$ ) due to the poor ohmic contact.

The capacitance voltage characteristics of the implanted subcollector DHBTs with Fe at  $I_c = 0$  mA, are plotted and compared with the standard mesa equivalent in Fig. 4.24 and those of the implanted subcollector DHBT without Fe from §4.4. From the figure, The implanted subcollector DHBTs with Fe exhibit a  $C_{cb}$  reduction of  $\approx 3$  fF over the entire measured range of bias voltages due to elimination of the charge dipole at the growth interface. At full depletion, this corresponds to a 25% decrease in  $C_{cb}$ , which corresponds to the area of the base access pad. In contrast, the implanted subcollector DHBT without Fe detailed in §4.4, exhibit the same reduction only at very high bias voltages. The standard, fully epitaxial mesa DHBTs have a larger collector base capacitance due to  $C_{cb,pad}$  being present.  $C_{cb,pad}$  is thus eliminated using the implanted subcollector DHBT process with Fe.  $A_{jBC} = 1.3 \times 5.25 \mu\text{m}^2$   $A_{pad} = 2.5 \mu\text{m}^2$ .  $C_{cb,pad}$  is thus  $\sim 28\%$  for this device.

The hybrid-pi model is extracted from the S-parameter and DC measurements. The S-parameters used to obtain the model correspond to those obtained at peak  $f_\tau$  and  $f_{max}$  which is at  $J_e = 6.8 \text{ mA}/\mu\text{m}^2$  and  $V_{cb} = 1.97$  V. The hybrid-pi model at  $I_c = 19.1$  mA and  $V_{cb} = 1.97$  V is shown in Fig. 4.44. The S -parameters of the actual device and the model are shown to closely agree in Fig. 4.45.

Device  $DC_7$  does not have the base pad isolated and its CV profile is shown in Fig. 4.46. The capacitance is the same for all three processes indicating that they are equivalent, as far as  $C_{cb,pad}$  is concerned. This is also further proof, that the decrease in capacitance is indeed due to the base access pad being isolated and not a fictitious

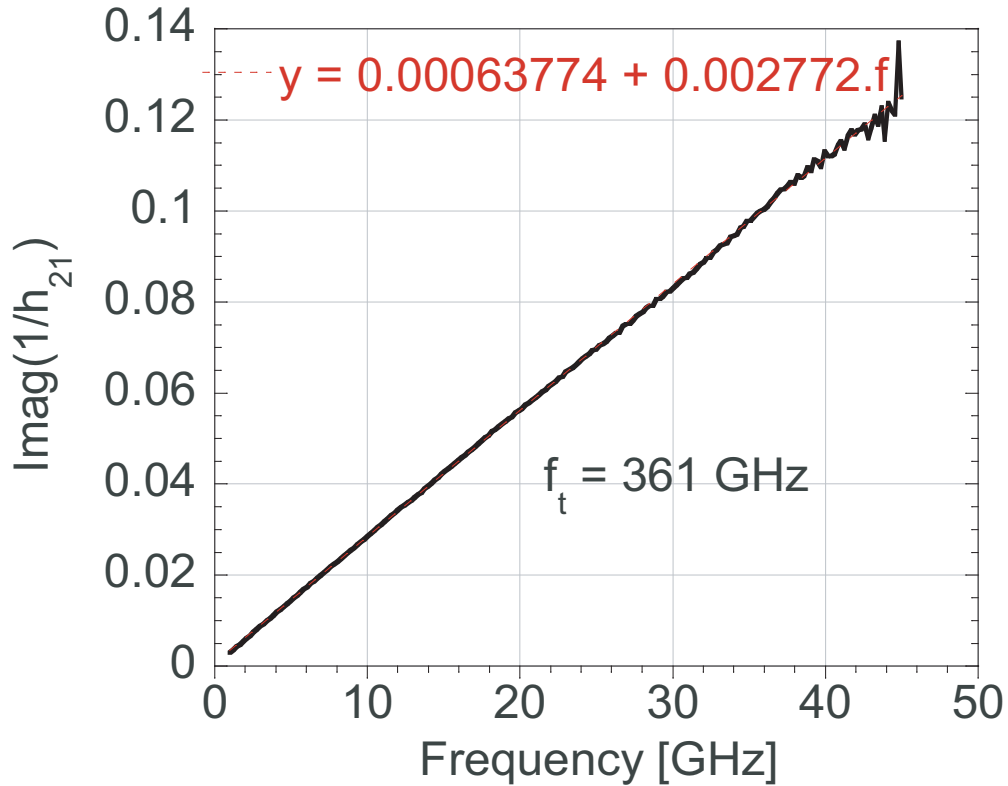


Figure 4.41: Extraction of  $f_\tau$ - The slope of  $\text{Imag}(1/h_{21})$  is  $f_\tau$

decrease due to variations in collector doping during growth, processing etc. For this device,  $A_{j,BC} = 2.1 \times 5.25 \mu\text{m}^2$ ,  $A_{pad} = 2 \mu\text{m}^2$ . This corresponds to  $\hat{C}_{cb} \approx 1.1 \text{ fF}/\mu\text{m}^2$ . Additional CV measurements of devices of different geometries shown in Fig. ?? determine the effective base collector area contributing to capacitance. Using  $\hat{C}_{cb} \approx 1 \text{ fF}/\mu\text{m}^2$  from Fig. 4.46, Table 4.8 shows the measured  $C_{cb}$  at full depletion for devices of various geometries and the effective pad area actually isolated, with subcollector implant boundary as drawn in Fig. 4.1.

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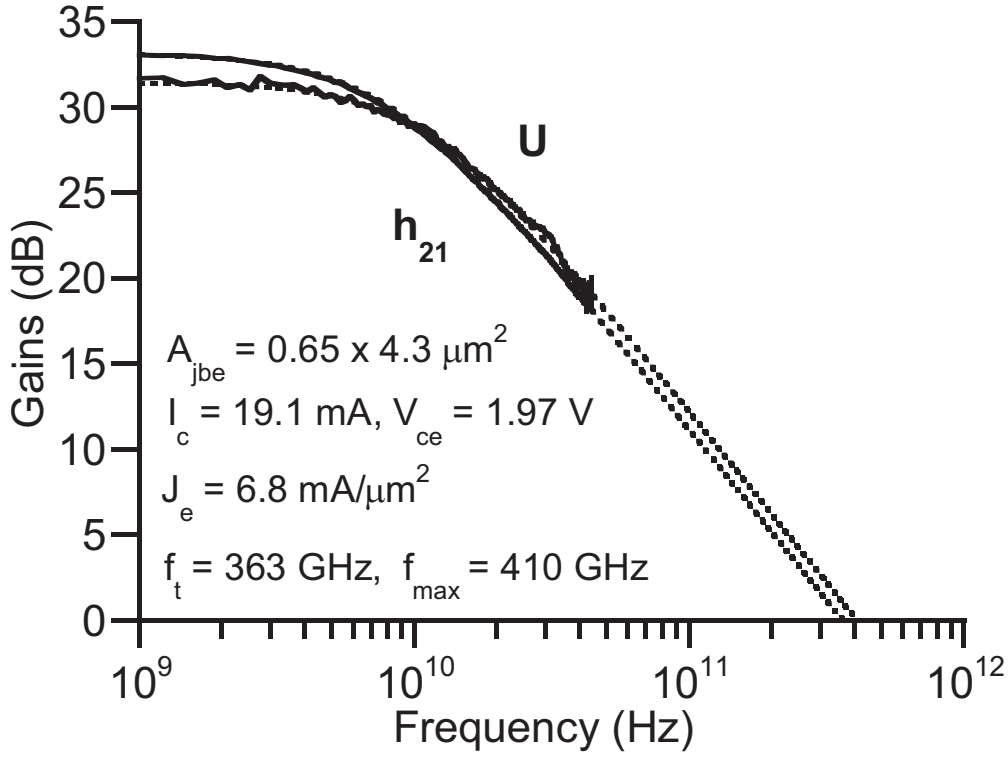


Figure 4.42: Microwave gains of Si implanted subcollector DHBTs

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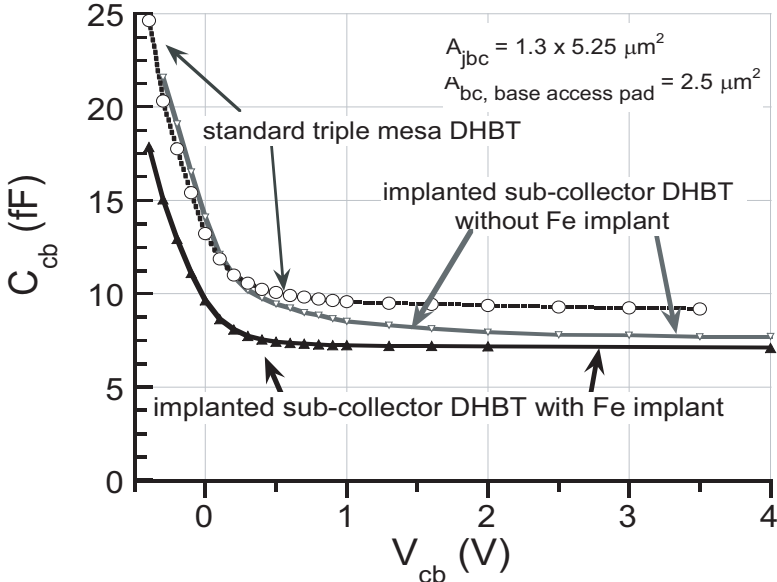


Figure 4.43: Capacitance voltage characteristics, at  $I_c = 0$  mA, of implanted subcollector DHBTs with the base pad isolated

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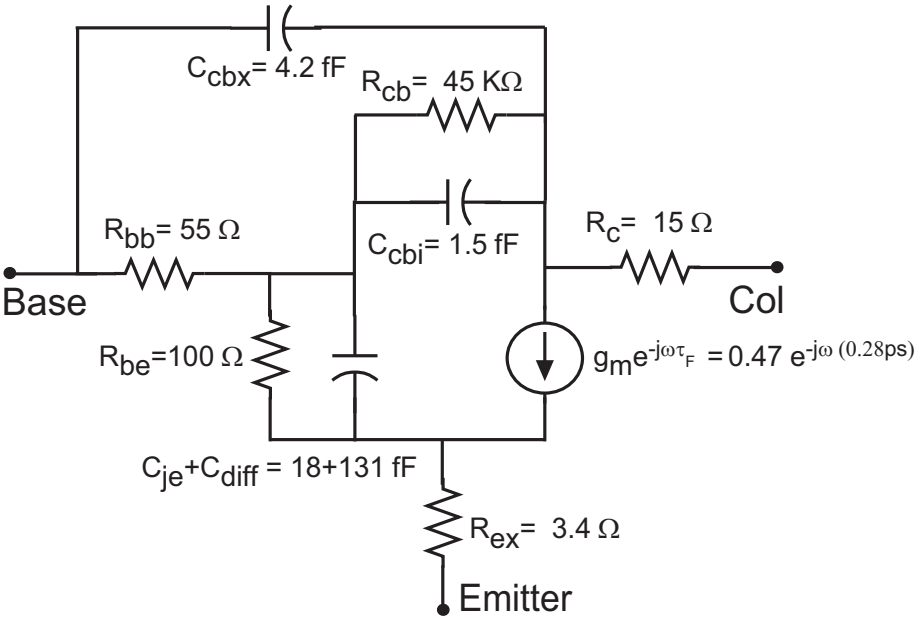


Figure 4.44: Hybrid- $\pi$  device model at peak  $f_\tau, f_{max}$

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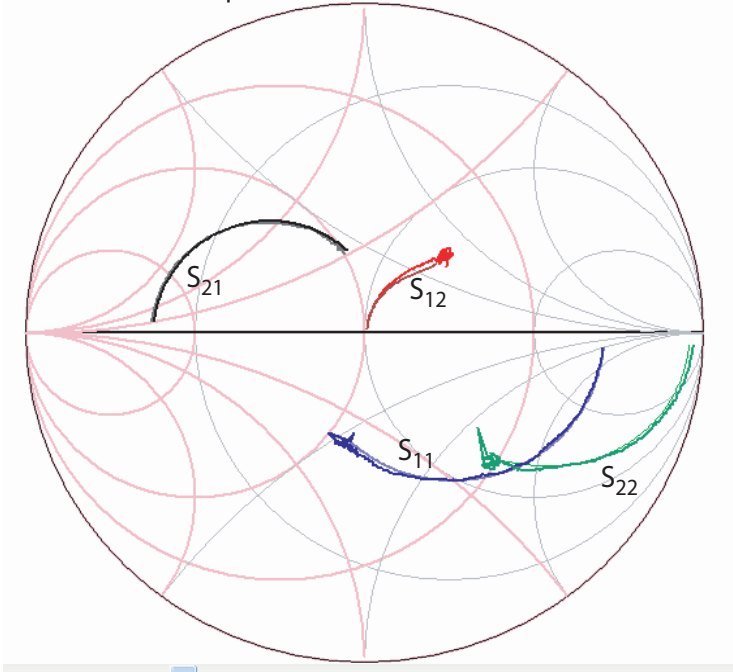


Figure 4.45: Measured S-parameters of the HBT and simulated S-parameters of the extracted hybrid- $\pi$  equivalent circuit

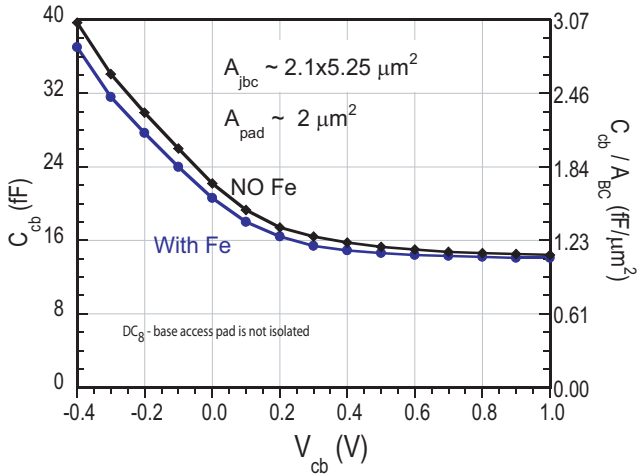


Figure 4.46: Capacitance voltage characteristics, at  $I_c = 0$  mA, of implanted subcollector DHBTS with the base pad *not isolated*

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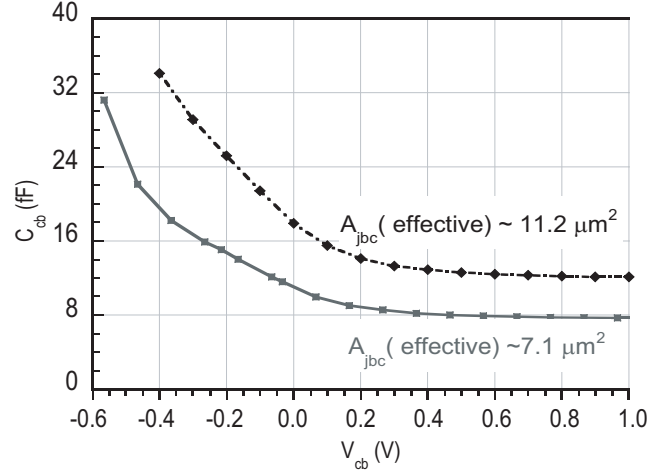


Figure 4.47: Capacitance voltage characteristics, at  $I_c = 0$  mA, of implanted subcollector DHBTs with Fe, of different geometries with the base pad *isolated*

The slightly lower  $C_{cb}$  reduction than expected, is due to longitudinal implant straggle. Thus  $C_{cb}$  in the extrinsic region can be reduced even further by optimizing the design of the subcollector implant boundary in the photo mask. The DC results compare with the standard mesa DHBTT and hence indicate the feasibility of manufacturing microwave HBTs on the doubly implanted substrates. The RF results indicate that  $C_{cb}$  is reduced corresponding to isolation of base access pad by implant, over the entire range of bias voltages. The interface charge dipole has been suppressed by the compensating Fe implant. The  $C_{cb}$  reduction would be more sig-

Table 4.8: Effective base access pad area isolated. The %  $C_{cb}$  reduction expected is in parenthesis

Device name	$A_{j,BC}$ on mask ( $\mu\text{m}^2$ )	$A_{pad}$ on mask ( $\mu\text{m}^2$ )	$C_{cb}$ at 1V (fF)	$A_{pad}$ isolated ( $\mu\text{m}^2$ )	Actual(expected) % reduction in total $C_{cb}$ (%)
DC1	$1.1 \times 5.25$	2.55	6.77	2.07	25(30)
DC3	$1.5 \times 5.25$	2.15	9.06	1.66	17(21)
DC4	$1.5 \times 7.25$	2.75	11.77	2.73	20(20)
DC5	$1.2 \times 5.25$	2.6	7.29	2.15	24(29)
DC6	$1.2 \times 7.25$	2.6	9.58	2.43	21(23)
DC7	$1.6 \times 7.25$	2.8	12.18	3.13	22(19)
DC15	$2.1 \times 5.25$	2.85	12.0	2.13	16(17)

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nificant for shorter length devices. The high collector contact resistance has resulted in lower  $f_\tau$  and  $f_{max}$  compared to [3]. The ohmic contact process can be fixed by reversing the order of the base and collector ohmics. This permits a high temperature anneal for the collector ohmics reducing contact resistivity to  $\approx 25 \Omega \cdot \mu\text{m}^2$ .



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# 5

## Implanted pedestal-subcollector DHBT

IN Chapter 4 , a technology is developed to eliminate the capacitance associated with the base access pad. However, lateral scaling of the collector base junction has not been achieved. In narrow mesa DHBTs shown in Fig. ?? in Chapter 2 ,

$$C_{cb} \propto \frac{W_{cb}}{V_{eb}} \quad (5.0.1)$$

$$R_{bb} \propto \coth \frac{L_T}{L_T} \quad (5.0.2)$$

where  $L_T$  is the ohmic transfer length limited by the ohmic contact to  $p^{++}$   $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}$ . Therefore any lateral scaling of the collector base area will result in a large increase in base resistance and this is the single biggest challenge facing InP mesa DHBTs today [1]. Furthermore, narrow base ohmic present a large access resistance and are difficult to form. As described in Chapter 2 , independent definition of the collector base junction becomes more important as HBTs are scaled. In this chapter, a novel idea is proposed and developed wherein implantation alone is used to independently scale the collector-base capacitance.

### 5.1 Various approaches to collector scaling

As described in Chapter 2  $C_{cb}$  scaling becomes important for future 250 nm HBTs. From Fig. ?? and Table 2.2, there is a large increase in  $f_{max}$ ,  $f_\tau$  and digital logic speed when the collectors are scaled.

The transferred substrate HBT in Fig. 5.1a dramatically scales the collector base capacitance. After base metal deposition, the wafer is planarized with BCB and then bonded to a carrier substrate. The wafer is then flipped over and the InP substrate is removed. A Schottky collector contact is then formed. Thus, the collector metal defines the base collector junction. This process has yielded devices with  $f_{max}$  of over a THz [2]. However, device yield was low due to the extreme complexity of the process which makes it unsuitable for even medium scale ICs.

## *CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT*

A popular approach is to greatly undercut the collector as shown in Fig. 5.1b. This cantilever like structure [4], is also susceptible to failures and is not used for high yield ICs.

A variant to the transferred substrate process is the buried electrode process[3] in Fig. 5.1c, where a W electrode defines the collector contact. The active devices are grown over this, so that the W contact defines the base collector junction. This, requires growth over the W metal which is complicated.

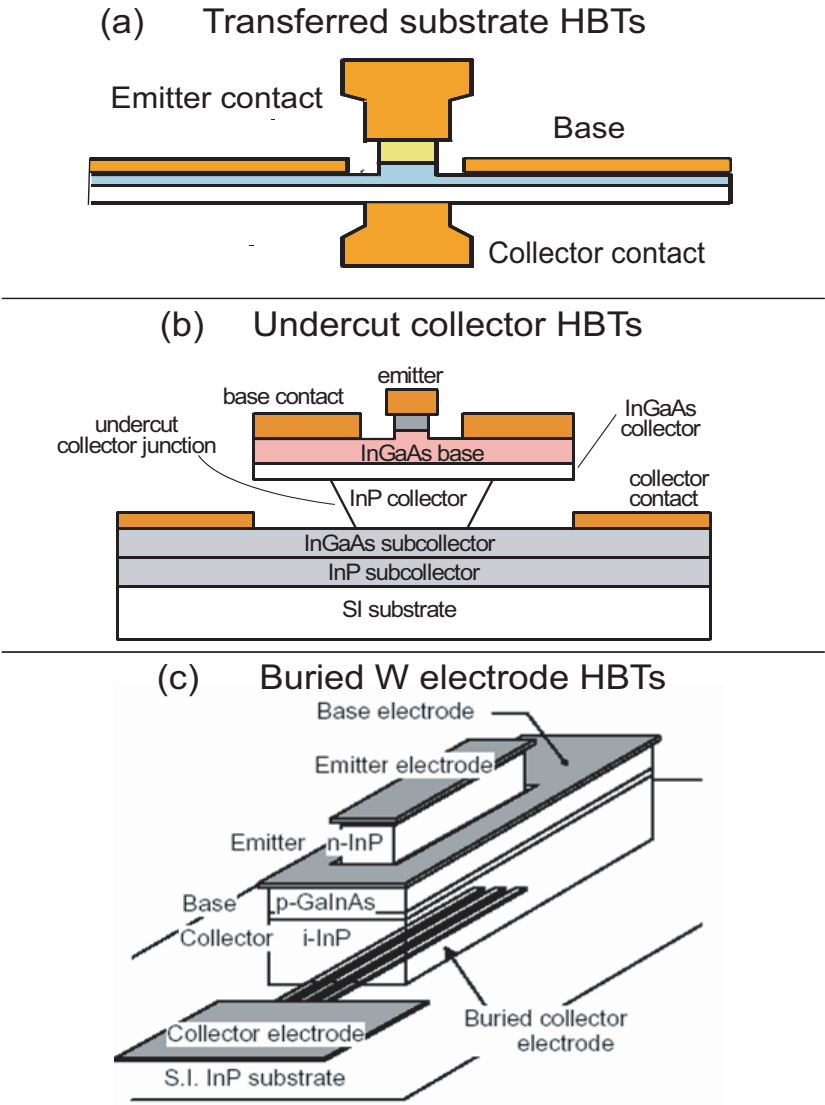


Figure 5.1: Various approaches to collector scaling

## CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

The narrow mesa HBTs manufactured today suffer from scaling challenges as described in Chapter 2 . Ion implantation offers the possibility of selectively defining junctions while maintaining a planar process and is exploited in modern SiGe HBTs [5]. One such process for collector scaling in InP HBTs, is the regrown collector pedestal process [6] whose device structure is shown in Fig. 5.2. The process starts with a template consisting of 200nm Undoped InP over 300 nm  $N^{++}$  InP sub-collector. Si is selectively implanted to form an  $N^+$  pedestal as shown in the figure. The drift collector, base, emitter layers are regrown and the device is formed. Thus, the intrinsic portion of the device over the collector has a collector depletion thickness defined by the thickness of the collector. In the extrinsic region, the depletion thickness is larger due to the undoped InP layer. Thus, the extrinsic capacitance is reduced, compared to mesa DHBT. The capacitances can be written as,

$$C_{cb,int} \propto \frac{W_p}{T_c} \quad (5.1.0)$$

$$C_{cb,ext} \propto \frac{W_{bc} - W_p}{T_c + T_p} \quad (5.1.0)$$

The regrown pedestal HBTs offer an alternative to the above complicated structures for collector scaling. However they [6] have the following drawbacks,

- DHBTs reported in [6] suffer from a process-dependent charge at the regrowth interface. A  $p^{++}$  InGaAs compensation layer is proposed as a solution but as seen in Chapter ?? , this is not repeatable
- Two MBE growths are required, the first forming the sub-collector and pedestal, and the second forming the active HBT layers
- The pedestal doping must be kept in the low  $1 \times 10^{18} \text{ cm}^{-3}$  range to minimize broadening of the pedestal through lateral straggle of the Si implant. This increases the access resistance in the pedestal and adds to the collector resistance
- These pedestal HBTs reduce, but do not eliminate the extrinsic base pad capacitance
- The collector contacts are offset from the collector by another 200 nm due to the undoped InP layer. This worsens the planarity compared to a regular mesa DHBT

An alternative idea is developed that addresses each of the above issues. This process provides an Fe implanted extrinsic semi-insulating layer for increased depletion depth in the extrinsic collector-base junction, a patterned buried subcollector

## CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

formed by a deep Si implant and a collector pedestal created by a second Si implant. In addition to the reduced extrinsic  $C_{cb}$ , and compensation of charge at the epitaxial growth interface, this provides the following enhancements: elimination of  $C_{cb,pad}$ , a single MBE growth, possibility of higher doping in the  $N^+$  pedestal, increased wafer planarity and hence potentially improved yield in the fabrication of large circuits.



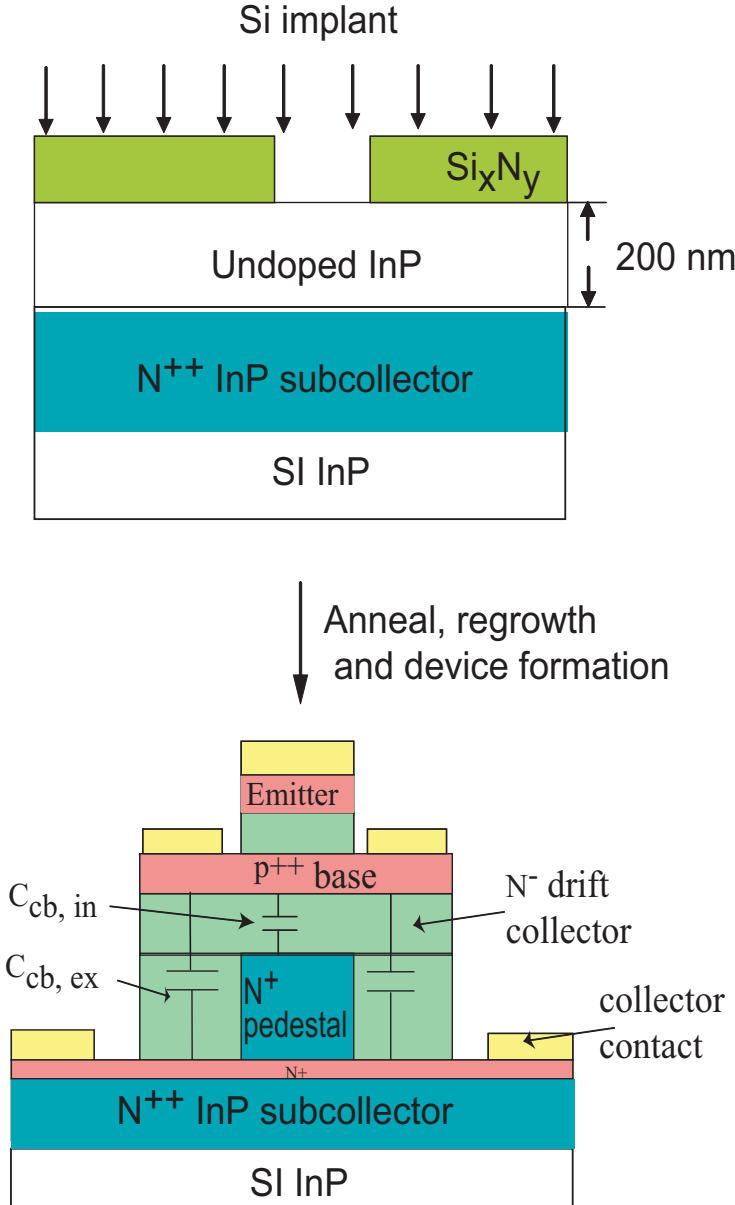


Figure 5.2: Pedestal HBT

## 5.2 Design of fully implanted pedestal-subcollector process

The process flow of the basic, implanted subcollector HBT is shown in Fig. 5.3. As before, alignment targets for stepper based lithography are defined on SI InP. The substrate is then implanted with Fe to a depth of  $\sim 0.2 \mu\text{m}$  at a doping of  $\sim 1 \times 10^{19} \text{ cm}^{-3}$  (Fig. 5.3a). The damage due to the implant is annealed out and the sample is then selectively implanted, as in Fig. 5.3b, with Si at a very high energy (350 keV) so that the peak (at  $\sim 0.5 \mu\text{m}$ ) lies well below the Fe implant peak. This forms a buried  $N^{++}$  subcollector. An  $N^{++}$  pedestal linking to the buried subcollector is then formed by a second patterned selective Si implant. Raised  $N^{++}$  pedestals for the collector contacts are simultaneously realized (Fig. 5.3c). The sample is then annealed at  $800 \text{ }^\circ\text{C}$  to activate the Si dopants. As before, the active InP DHBT layers, the  $N^-$  drift collector, base, and emitter layers are grown by MBE. Devices are formed by wet-etching the emitter and base mesas. HBT isolation does not require a subcollector mesa etch decreasing the total HBT mesa height by  $500 \text{ nm}$ . There are several advantages,

- The Fe implant forms a SI layer, equivalent to the undoped InP layer in the regrown pedestal HBT, over the  $N^{++}$  subcollector. The  $N^{++}$  subcollector is also formed by ion implantation. Thus this fully implanted pedestal-subcollector process requires no additional regrowth.
- The Fe implant automatically compensates the interface charge over a very wide range of charge densities
- The buried subcollector does not extend beyond the length of the emitter, similar to the implanted subcollector DHBTs with Fe. Thus the capacitance due to the base access pad is eliminated.
- The high Fe doping in the SI layer  $\sim 5\text{-}8 \times 10^{18}$  means that a high concentration of Si in the pedestal is permissible, since the straggle is now defined by the lateral distance at which the pedestal doping falls below the background Fe in the SI region.
- Raised  $N^{++}$  pedestals are simultaneously realized for collector contacts. Device isolation etch is not required due to the buried, isolated  $N^{++}$  subcollector. Thus the device planarity is better than a triple mesa DHBT by  $\sim 500 \text{ nm}$
- It is very modular and can be inserted into any modern InP mesa HBT process.

CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

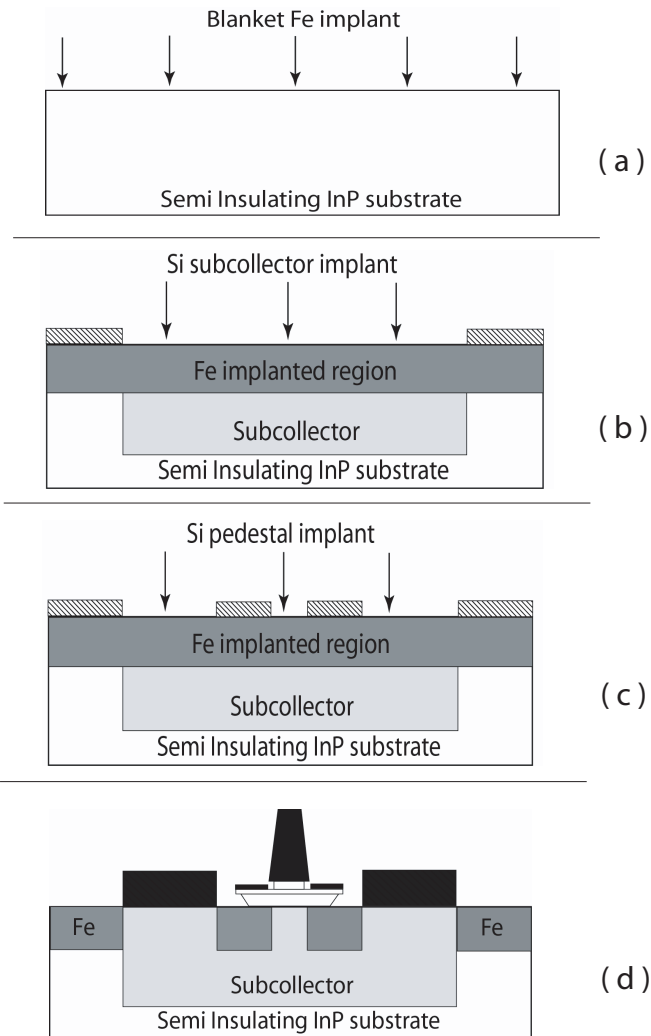


Figure 5.3: Implanted pedestal-subcollector DHBT process

## CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

Thus the fully implanted process, is an alternative to the regrown pedestal HBT and provides several enhancements. The various challenges of the implanted pedestal-subcollector process are,

- Low sheet resistance subcollector
- Formation of thick SI Fe implanted region
- Formation of  $N^{++}$  pedestals with low resistivity and low lateral straggle
- crystalline, defect free semiconductor before growth
- excellent surface morphology, comparable to InP substrate, prior to growth
- defect free MBE growth  $\Rightarrow$  DC characteristics similar to epitaxially grown HBTs

Each issue is addressed in the subsequent sections.

### 5.2.1 Formation of buried $N^{++}$ subcollector

The Si implant forms the subcollector as in the implant subcollector process. However, unlike the implanted subcollector DHBT, the main idea behind the implanted pedestal-subcollector DHBT is a deep Fe implant since it is used to compensate  $\approx$  the top  $0.2 \mu\text{m}$  of the subcollector. As seen in Chapter 3, it is not easy to compensate very high  $N^{++}$  doping levels. Therefore, the Si implants should be of very high energy so that the peak lies well below  $0.2 \mu\text{m}$ . This ensures that the Gaussian distribution (of the implanted Si) tails to  $\sim 5 \times 10^{18} \text{ cm}^{-3}$ , doping level at which Fe is reported to successfully compensate the  $N^+$  doping. The highest commercially available Si doping at  $200^\circ \text{C}$  is 350 keV. Further, it is seen that increasing the thickness of the SI region, is not beneficial as might be expected. The lateral broadening due to implant straggle is, to first order,  $\sim T_p$ , the height of the  $N^+$  pedestal. If  $T_p$  is increased,  $C_{cb,ext}$  decreases due to larger extrinsic depletion depth, but  $C_{cb,int}$  increases due to a wider effective  $N^+$  pedestal. Thus the two effects offset each other and for the minimum, optically resolvable implant mask dimension at UCSB of  $\sim 0.5 \mu\text{m}$ , it is seen that  $C_{cb}$  decrease is not significant beyond  $T_p$  of  $0.2 \mu\text{m}$ .

Fig. 5.4 shows the SIMS profile and the simulated TRIM distribution for the implant conditions in Table 5.1 where Si and Fe are co-implanted. There is no diffusion of the Fe and Si as implanted. However, the fluences were slightly off, due to the poor calibration of the implant system. (This is not an artifact due to charge-mass interference in SIMS since  $^{29}\text{Si}$  and  $^{54}\text{Fe}$  also yielded similar results).

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Table 5.1: Implant conditions for implanted pedestal-subcollector HBT process

Implant species	Implant energy	Implant fluence	Offset angle	Implant temperature
Si	350 KeV	$5 \times 10^{14}$ ions/cm <sup>2</sup>	7 °	200 ° C
Fe	150 KeV	$2.4 \times 10^{14}$ ions/cm <sup>2</sup>	7 °	200 ° C
$I_{ions} \leq 50 \mu A$				

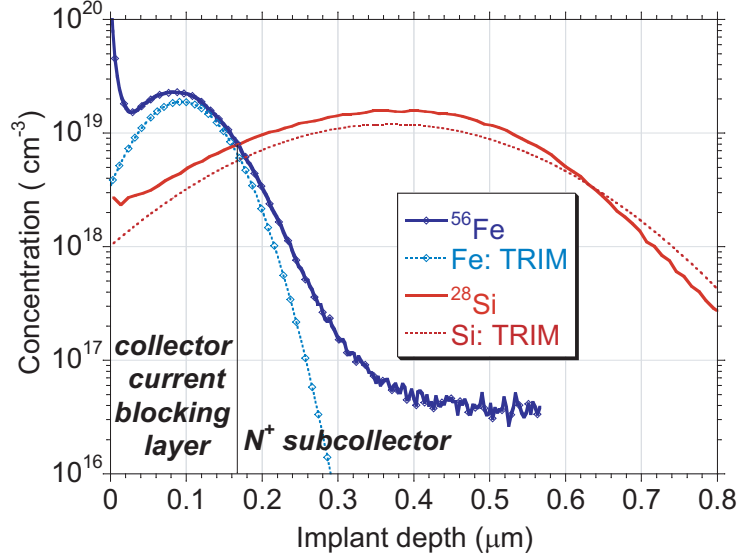


Figure 5.4: Distribution of Fe and Si from SIMS and simulated by TRIM

The shallow Fe implant in Chapter 4 is annealed at 700 °C for 5 min. The same anneal conditions for Fe are employed here. This is shown in §5.2.2 to anneal the implant damage and activate the dopants. Fig. 5.5 shows the profile after such an anneal. There is significant depletion of Fe atoms and a peak of Fe is formed at  $\sim 0.14 \mu\text{m}$  which is  $R_p + \Delta R_p$ . This is well observed in literature and is described in Chapter 3. As seen from the plots, the Fe concentration falls below the Si concentration at a depth of  $\sim 0.16 \mu\text{m}$ . Assuming 100% activation of Si and Fe,  $T_p \sim 0.16 \mu\text{m}$ . While  $T_p = 0.2 \mu\text{m}$  is desirable, these were nevertheless chosen for the first generation implanted pedestal-subcollector HBTs.

For these implant conditions, the sheet resistance is determined from Hall measurements and is shown in Table 5.2. The negative Hall voltage indicates that the

Table 5.2: Hall measurements of resistivity and mobility for deep Si and Fe implant

Anneal temperature (°C)	Anneal time (s)	Hall mobility ( $\text{cm}^2/\text{V} \cdot \text{s}$ )	Carrier density ( $\text{cm}^{-2}$ )	Sheet resistance ( $\Omega/\square$ )
700	300	-1194	$-2.0 \times 10^{14}$	25.8
700	600	-1224	$-1.9 \times 10^{14}$	26.2
725	600	-1189	$-2.0 \times 10^{14}$	25.7
750	300	-1236	$-2.1 \times 10^{14}$	24.5

carriers are electrons. The sheet resistance is nearly the same for all the various an-

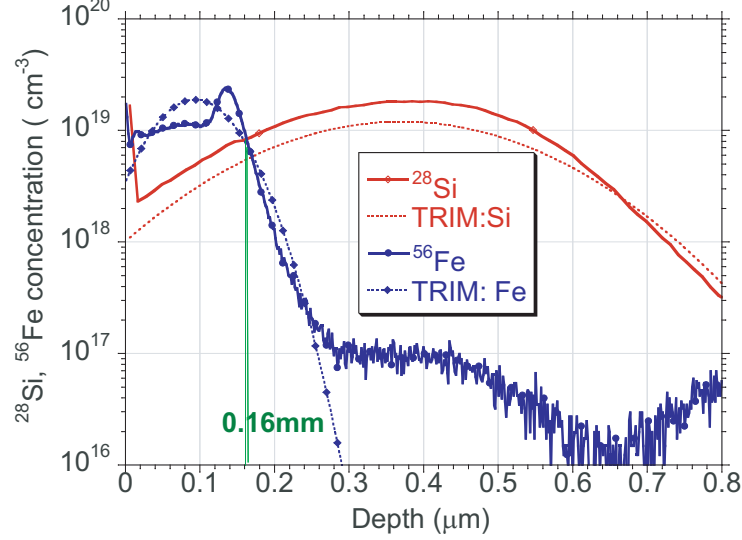


Figure 5.5: Distribution of Fe and Si after anneal

neals. However, 700 °C for 5 min gave rise to minimum defects due to the anneal. From the above data and measured concentration profiles, the activation of Fe and Si can be calculated. Assuming that the active Fe compensates Si upto 1632 Å, and integrating the Si concentration from this point,  $n_{Si} = 6.5 \times 10^{14} \text{ cm}^{-2}$ . From Hall measurements,  $n_{Si,eff} = 2.5 \times 10^{14} \text{ cm}^{-2}$ . Therefore activation of Si is  $\sim 39\%$ . The active Si concentration at 1632 Å is therefore  $3.8 \times 10^{14} \text{ cm}^{-3}$ . Since one Fe atom compensates one Si atom, minimum activation of Fe is  $N_{Fe}/N_{activeSi}$  at 1632 Å = 41 %.

## 5.2.2 Crystallinity

It is seen from Chapter 3 that  $^{56}\text{Fe}$  causes significant lattice damage. The crystallinity is evaluated from Ion channeling(RBS) measurements. The defects are quantified in terms of displaced  $In$  atoms. If displaced  $In$  fraction is 1, it is completely amorphous and if it is  $\leq 0.02$ , it is crystalline. Sample is completely amorphous in the top 1500-2000 Å suggesting that Fe is creating most of the damage and that defects due to Si is much smaller. The  $In$  concentration in  $InP$  is  $5.3 \times 10^{22} \text{ atoms-cm}^{-3}$  and from the displaced  $In$  fraction in Fig. 5.7, the total number of  $In$  atoms displaced can be determined. From the discussion on Fe implants in Chapter 3, it is seen that prolonged anneals at temperatures of  $\sim 700 - 750$  °C are required for damage removal and activation. Anneals at over 725 °C for prolonged periods

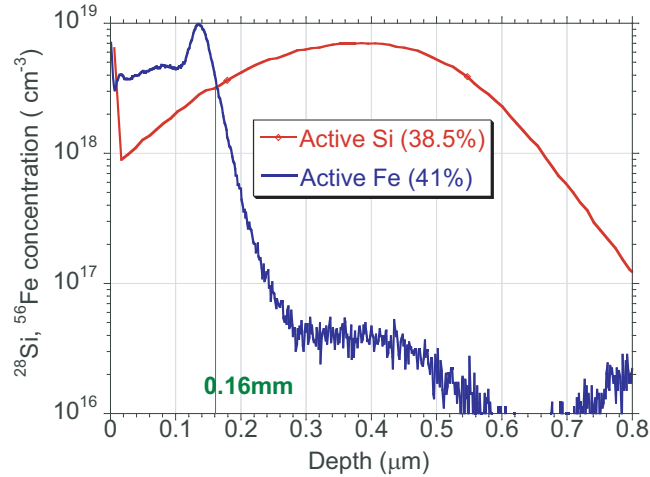


Figure 5.6: Distribution of Active Fe and Si calculated from Hall and SIMS

gave rise to several In rich zones due to failure of the  $\text{Si}_x\text{N}_y$  cap. The samples are annealed at  $700^\circ\text{C}$  for 5 min. and the crystallinity is checked using RBS. As seen in Fig. 5.8 and Fig. 5.9, samples implanted with lower doses ( $\leq 2.4 \times 10^{14}$  ions/ $\text{cm}^2$ ) of Iron are completely recovered. There is surface damage where the implant fluence of Iron is higher ( $1 \times 10^{15}$  ions/ $\text{cm}^2$ ). The uncertainty for In defect measurement is 1 to 2 %.



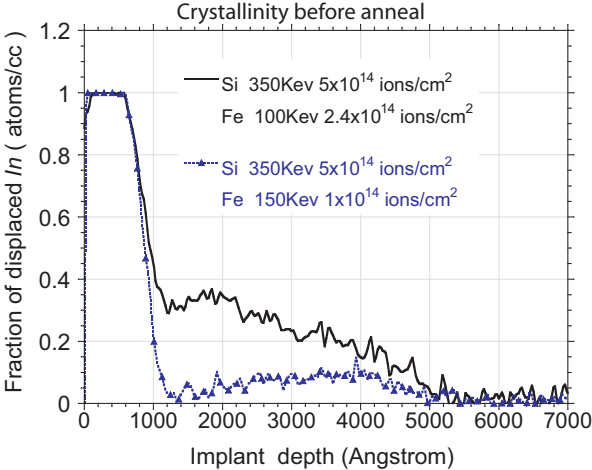


Figure 5.7: Crystallinity with co-implants of Si and Fe before any anneal

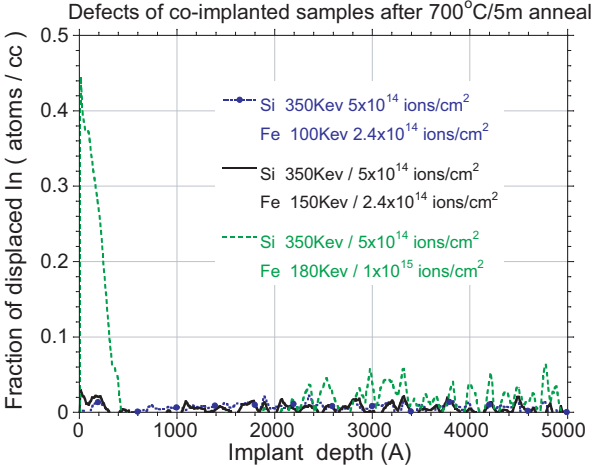


Figure 5.8: Crystallinity of Si and Fe co-implanted InP annealed at 700 °C/5 min

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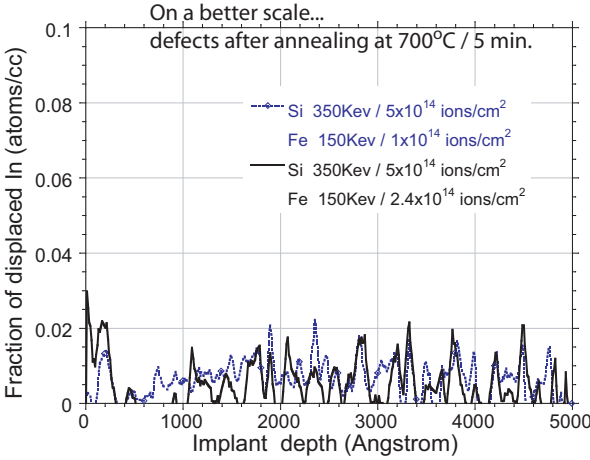


Figure 5.9: ...A more convincing scale

### 5.2.3 Surface morphology

Fig. 5.10 show the surface scans for various conditions. As implanted with Si and Fe, the surface is quite rough,  $\sim 18$  nm. When annealed at  $800^\circ\text{C}$  for 10 sec. the surface roughness reduces to  $\sim 12$  nm. As seen from the previous section and from §3.1.3 in Chapter 3 , heavy acceptor elements in InP generally require prolonged anneals at temperatures over  $650^\circ\text{C}$ . AFM scans of this co-implanted substrate, annealed at  $700^\circ\text{C}$  for 5 min. indicates that the surface quality is comparable to virgin SI InP in Fig. ?? in Chapter 3 . This further validates the use of these annealing conditions for Fe implant.

### 5.2.4 The Fe implanted extrinsic layer

The Fe layer serves to increase the depletion depth in the extrinsic portion of the device under the base contact. Co-implantation of Si and Fe causes a difference in activation as opposed to the case when

- Fe is implanted first and annealed
- Si is then implanted and annealed

When Fe is implanted first and activated they have occupy *In* sites. The Si therefore has less sites to occupy.

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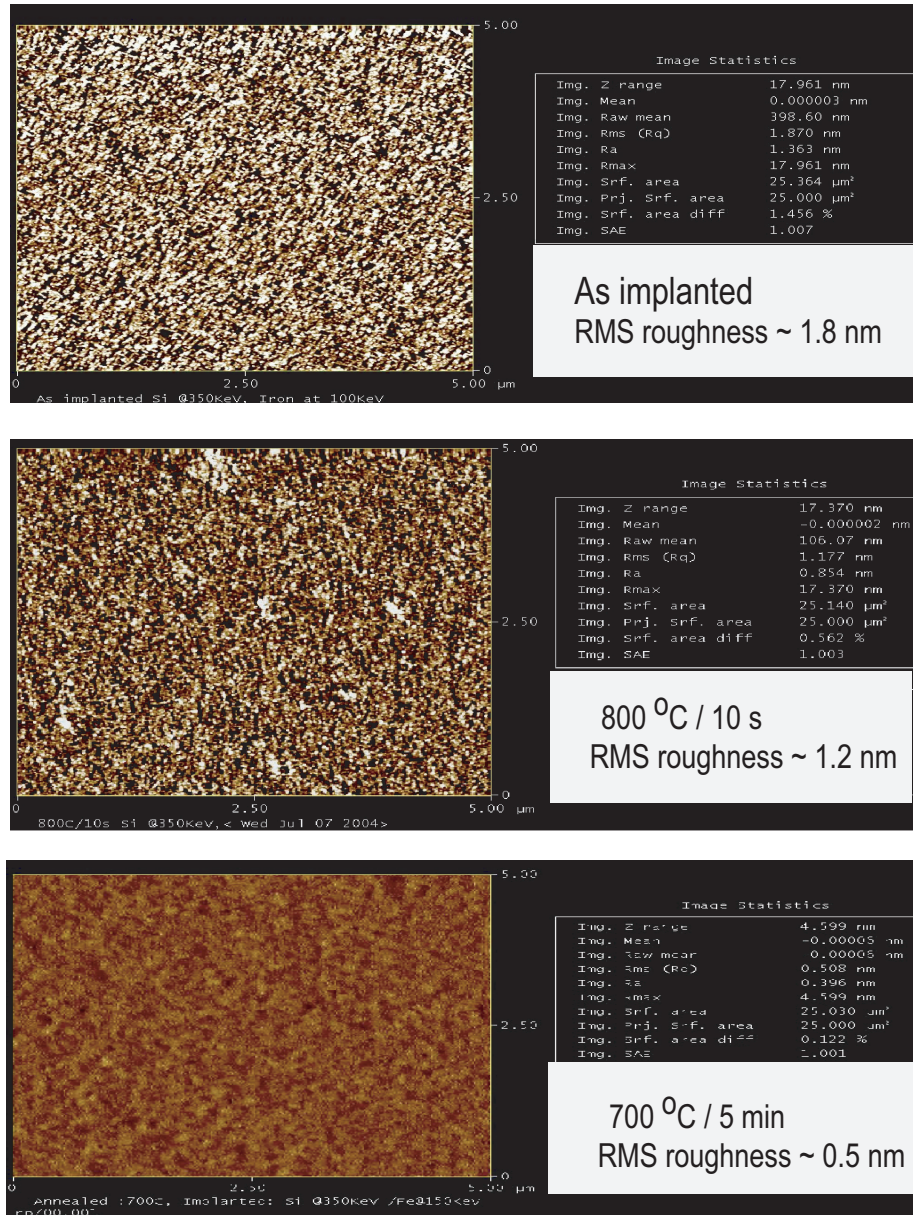


Figure 5.10: Surface scans of the Si and Fe co-implanted substrate

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The current flow mechanism of an  $N^{++}$ -SI- $N^{++}$  and  $p^{++}$ -SI- $N^{++}$  layer stacks respectively is developed in [7]. The main purpose of the SI region is to decrease the capacitance in the extrinsic region by increasing the distance between the two charge dipoles in the base and subcollector. C-V measurements are used to check the presence of an active SI layer and if present, the thickness of this region. For this purpose, Schottky diodes are formed. The samples are non-selectively implanted first with Fe at the implant conditions, annealed at 700 °C for 5 min. They are then non-selectively implanted with Si and annealed at 800 °C for 30 sec. The implant conditions are the same as in §5.2.1. Schottky contacts are placed on the top SI layer, while ohmic contacts are placed on the  $N^{++}$  subcollector. Fig. 5.12 shows the schematic of such a structure.

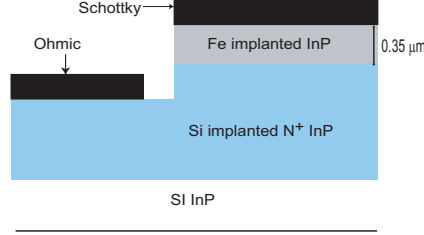


Figure 5.11: Schottky diodes on Fe and Si implanted substrate

The contacts placed on the  $N^{++}$  regions are verified to be ohmic by checking the I-V characteristics (Fig. 5.13). Similarly the contacts on the top surface, implanted with Fe are verified to be Schottky by measuring the I-V characteristics of this SI- $N^{++}$  diode. Fig. 5.14 also shows the Schottky diode characteristics of such a structure on a fully epitaxial structure ( $0.2 \mu\text{m UID} / 0.3 \mu\text{m } N^{++} \text{ InP}$ ), which is the starting template in a regrown pedestal HBT process. The forward characteristics of the Schottky diode on the fully implanted structure indicates a very high *turn on* voltage. From the theory of transport in a SI region, the current increases sharply once the Trap Filled Voltage Limit is reached. From Fig. 5.14,  $V_{TFL}$  is  $\sim 2 \text{ V}$ . In the case of a Schottky diode on the epitaxial template, the theory of transport is similar to a forward biased Schottky junction and the current sharply increases at  $\sim 0.3 \text{ V}$ . In the reverse bias case, the situation is quite different. The depletion region thickness at zero bias for the Schottky diodes on  $0.2 \mu\text{m UID} / 0.3 \mu\text{m } N^{++} \text{ InP}$  is  $\approx$  thickness of the UID region =  $0.2 \mu\text{m}$ . The reverse diode current characteristics are dominated by the thermal generation in the intrinsic region. The breakdown is expected to be large in this case, due to the wideband gap of InP and the large depletion thickness. The thickness of the space charge region for the SI- $N^{++}$  junction is calculated from the active Fe and Si doping profiles obtained in the previous case.

$$x_A = \sqrt{\frac{2\phi_{bi}}{qN_A}} \quad (5.2.0)$$

where,  $N_A$  is the active Fe concentration in the SI region,  $\phi_{bi}$  is the built-in potential of the SI- $N^{++}$  junction. Since  $E_{Fp}$  of the SI region  $\sim E_A$  and  $E_{Fn}$  of  $N^{++} \text{ InP} \sim E_c$ ,  $\phi_{bi} \sim 0.7 \text{ eV}$ . Assuming uniform  $N_A \sim 1 \times 10^{18} \text{ cm}^{-3}$  and  $N_D \sim 1 \times 10^{19} \text{ cm}^{-3}$ ,  $x_A = 88 \text{ \AA}$ . Due to the small depletion region, the breakdown mechanism is possibly dominated by tunneling currents which could explain the dramatic increase in current at  $\sim 1.5 \text{ V}$ . The C-V characteristics in Fig. 5.15 of such the Schottky diodes indicate that there is a thick SI region. From the capacitance at zero bias,

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the thickness of the SI region is deduced to be  $\sim 177$  nm.  $x_{dep} \sim 200$  nm for the Schottky diode on the epitaxial UID InP/  $N^{++}$  InP, as expected since the thickness of the UID InP region as grown is 200 nm. The capacitance is fairly flat with voltage indicating that beyond the depletion thickness, the region is fairly  $N^{++}$ , such that the depletion region does not shift much with bias. Fig. 5.16 compares the C-V characteristics of two cases

- Fe is implanted and activated first followed by Si implant and activation
- Fe and Si are co-implanted and activated together

The depletion thickness is lower in the case when Si and Fe are co-implanted and activated together. This is possibly due to Si and Fe competing for the same In sites between 160 - 180 nm when Si and Fe concentrations are comparable, resulting in lower active Fe concentration, decreasing the thickness of the SI region.

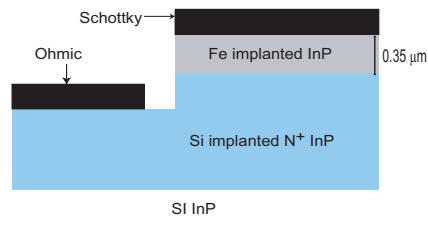


Figure 5.12: Schottky diodes on Fe and Si implanted substrate

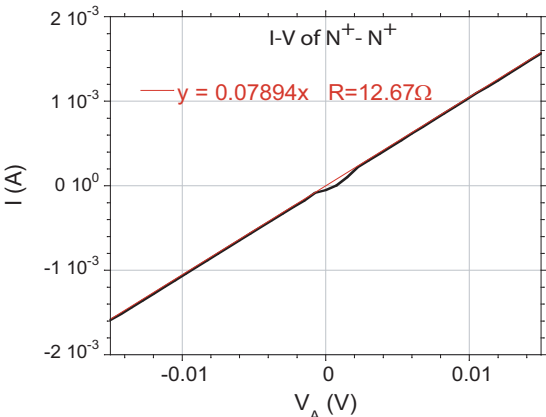


Figure 5.13: Ohmic transport between  $N^{++}$  pads

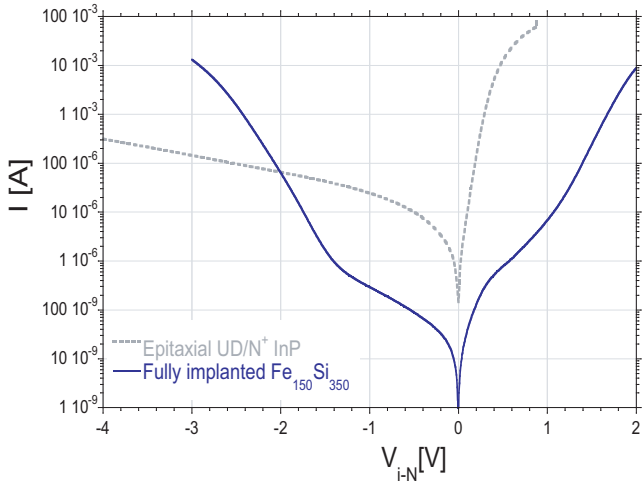


Figure 5.14: Sckottky diode characteristics of Fe and Si implanted substrate



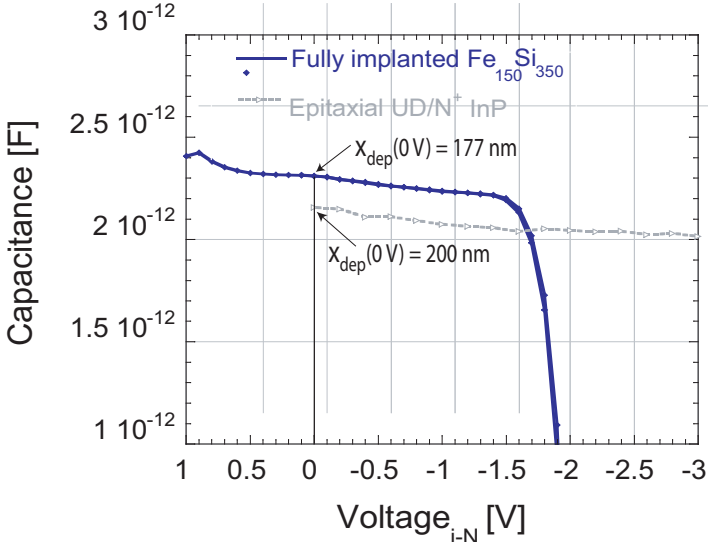


Figure 5.15: C-V of the Schottky diodes on Fe and Si implanted substrate, indicating the thickness of the SI depletion region

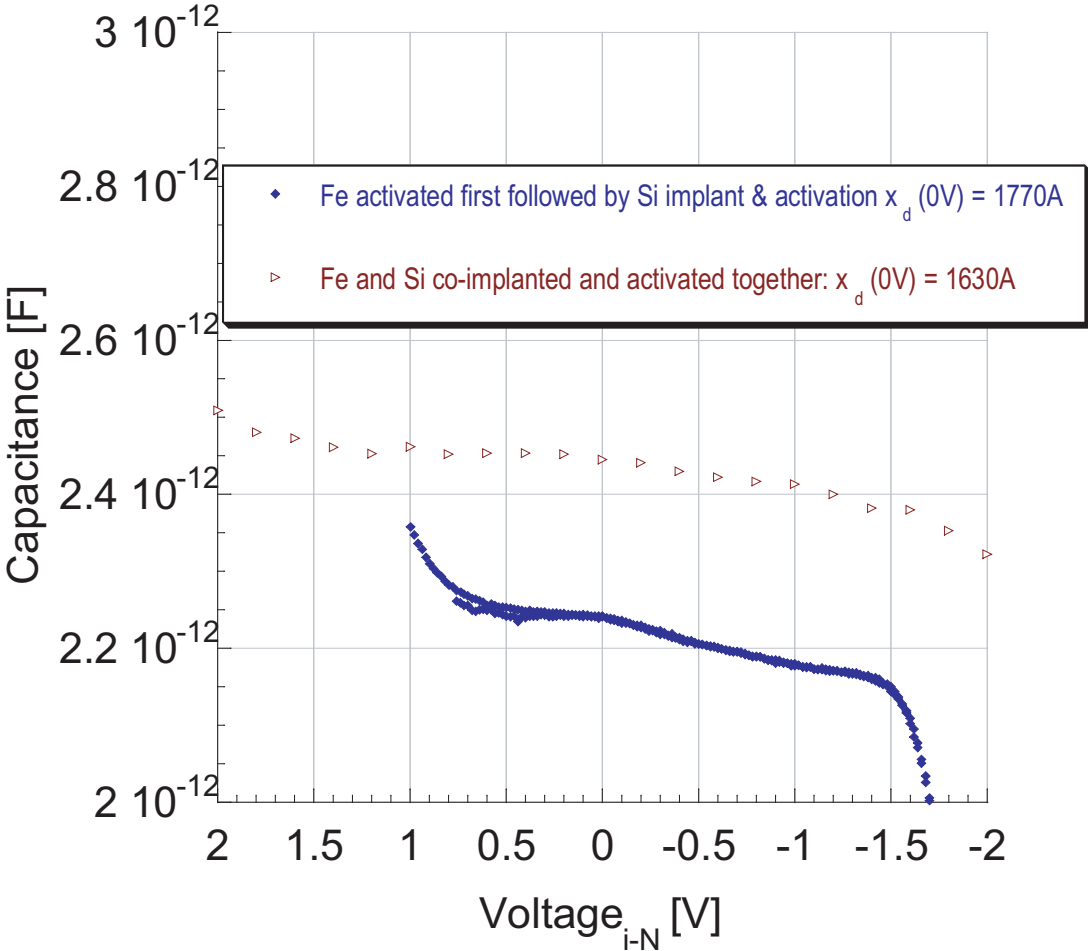


Figure 5.16: C-V of the Schottky diodes when Fe and Si are co-implanted and activated together, and when Fe is activated first followed by Si

CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

Table 5.3: Implant conditions for formation of  $N^{++}$  pedestal

Implant species	Implant energy	Implant fluence	Offset angle	Implant temperature
Si	100 KeV	$5 \times 10^{14}$ ions/cm <sup>2</sup>	7 °	200 ° C
	30 KeV	$8 \times 10^{13}$ ions/cm <sup>2</sup>		
	10 KeV	$4 \times 10^{13}$ ions/cm <sup>2</sup>		
$I_{ions} \leq 10 \mu A$				

### 5.2.5 Formation of $N^{++}$ pedestal

In §5.2.1, the active Fe and Si concentrations are computed. The pedestal doping needs to overwhelm the Fe concentration. Si is implanted at the conditions given in Table 5.3 in a substrate where Fe has been implanted and activated.

After annealing at 800 °C for 30 sec., Hall measurements indicate that the sheet resistance is  $\sim 38 \Omega/\square$  and for  $T_p \sim 180$  nm, resistivity of the pedestal is  $\sim 6.5 \Omega \cdot \mu m$ . When both the pedestal and subcollector are present, the sheet resistance of the combined stack is  $\sim 17 \Omega/\square$  and  $n_{Si} = 4.7 \times 10^{14} \text{ cm}^{-2}$  and the mobility is  $800 \text{ cm}^2/V \cdot s$ . Fig. 5.17 shows the concentration of Si and Fe after all three implants and anneals. With the chosen implant conditions, Si overwhelms the Fe peak.

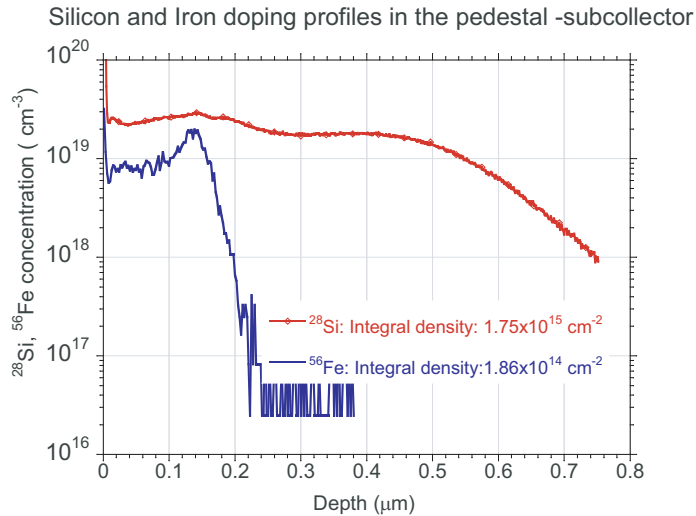


Figure 5.17: Concentration of Si and Fe after all three implants

It is seen that an anneal of 700 °C for 5 min. is required for Fe activation and damage Si removal. Good activation of Si is obtained only after annealing at 800 °C.

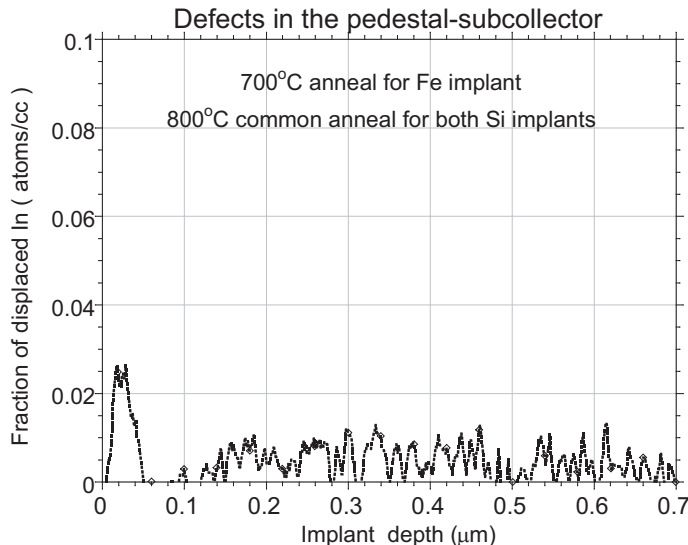


Figure 5.18: Crystallinity after all implants and dry etch

Since there are two Si implants, one for the subcollector and one for the pedestal, the question arises if two separate anneals are required to activate the Si dopants. It is seen that the TiW alignment marks does not hold up to two anneals at 800 °C. Fig. 5.18 shows the defect density after all three implants. The implant conditions for the Fe and Si subcollector implant are the same as in §5.2.1 and the above conditions are used for the Si pedestal implant. After the Fe implant, the sample is annealed at 700 °C for 5 min. Si is implanted for the subcollector and then the Si pedestal implant is carried out. Both the implants are annealed together at 800 °C for 30 sec. Fig. 5.18 shows that the defect density is within the noise of the measurements, indicating crystallinity of the InP substrate after 3 implants and 1 anneal for Fe and 1 common anneal for Si. Since these are identical implants, at different depths, a single anneal does not yield significantly different results for resistivities for the various layers. Fig. 5.18 shows the defect density after implants and also 3 dry etches; for the alignment targets, subcollector implant window, and pedestal implant window.

### Lateral straggle due to the pedestal implant

As defined in Chapter 3 , the straggle is defined as the point when the active, lateral Si concentration falls below the background Fe doping. In the case of SI InP substrate, the active Fe doping is between  $5 \times 10^{16} \text{ cm}^{-3}$  and  $9 \times 10^{16} \text{ cm}^{-3}$ .

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Therefore the lateral straggle is appreciable. The thick SI region over the  $N^{++}$  subcollector is formed by Fe implant, and has an active Fe concentration of  $5 \times 10^{18} \text{ cm}^{-3}$ . Fig. ?? is a plot of the lateral straggle calculated from *TRIM*. The pedestal doping is seen from Fig. 5.17 to be  $\sim 2 \times 10^{19} \text{ cm}^{-3}$  and uniform Fe doping of  $5 \times 10^{18} \text{ cm}^{-3}$  is assumed. The lateral spread is  $\sim 0.12 \mu\text{m}$  when the Si concentration falls below the active Fe doping assumed to be  $\sim 5 \times 10^{18} \text{ cm}^{-3}$ . This is not an exact solution since the Fe doping itself varies with distance. The exact calculation can be performed; however, *TRIM* profile of Fe implant has to be relied upon. From §5.2.1, the *TRIM* calculation is not representative of the Fe profile due to the diffusion of Fe during anneal. For this calculation the *TRIM* profile of Fe is assumed, with 40% activation of these Fe acceptors. In this case, the lateral straggle can be calculated as the lateral distance( $z$ ) at which

$$n_{Si}(x, z) \exp(-z^2/2\pi\Delta R_z^2) = n_{Fe}(x, z) \quad (5.2.0)$$

where  $\Delta R_z$  is the lateral spread of the 100 KeV Si implant calculated from *TRIM* to be  $577 \text{ \AA}$ . The lower energy implants are assumed to contribute insignificantly to this straggle. Fig. 5.20 shows the lateral straggle distance  $z$  which is  $\sim 0.25 \mu\text{m}$ .

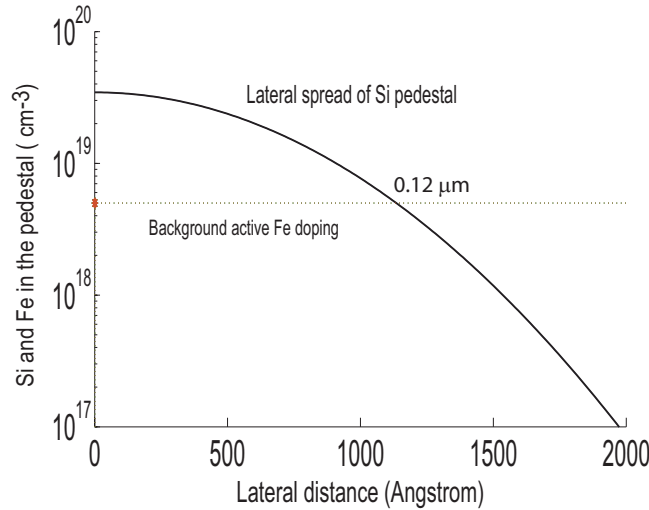


Figure 5.19: Lateral spread in the pedestal due to implant, assuming  $2 \times 10^{19} \text{ cm}^{-3}$  doping at the mask edge. The lateral straggle is defined as the point when the Si concentration falls below the active Fe doping  $\sim 5 \times 10^{18}$

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The resistive straggle can be measured using special *dog-bone* TLM structures shown in Fig. 5.21. The spacing between the ohmic regions and the width of the

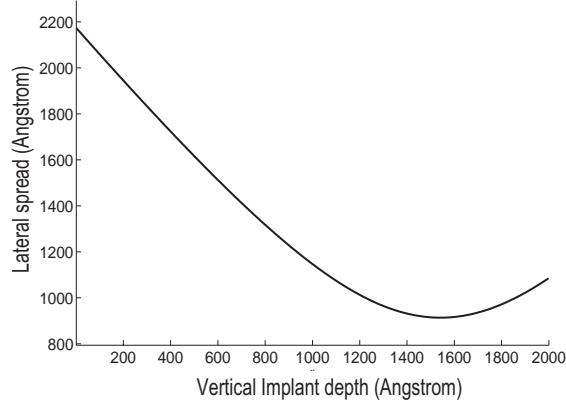


Figure 5.20: Lateral spread in the pedestal due to 100 KeV Si implant. The lateral straggle is defined as the point when the Si concentration falls the active Fe doping calculated from *TRIM* with Fe activation assumed to be 40 %

TLMs are varied. The width of the TLM should be comparable to the straggle, which renders standard TLMs for measuring contact and sheet resistivities useless for this purpose. The wafer is first implanted with Fe and annealed and then the pedestal Si implant is carried out selectively in the shaded regions shown in Fig. 5.21. Standard TLMs can measure, the sheet resistance and contact resistivity. The measured resistance for each TLM (of widths  $w_1, w_2 \dots$ ) is plotted vs. the TLM spacing  $x$  and the slope  $m_{str}$  is calculated. The lateral resistive straggle is then calculated as follows

$$R_{sh} = m_{str} \cdot (w + \Delta w) \quad (5.2.0)$$

where  $w$  is the width of the TLM. Plotting  $\frac{R_{sh}}{m_{str}}$  vs. the variation in TLM width as drawn on the mask  $w$ ,  $\Delta w$  can be obtained. This is the lateral resistive straggle shown in Fig. 5.22. Note, the resistive lateral straggle also includes the effects of widening of the  $Si_xN_y$  implant mask during processing. Fig. 5.22 shows the resistive straggle of the Si pedestal implanted in an already Fe implanted substrate. The resistive straggle,  $\Delta w \sim 0.1 \mu m$ , which means that the implant widens by 50 nm on either side of the implant mask. However, it should be pointed out that resistive straggle is a pessimistic calculation and the capacitive straggle is more relevant. When the doping falls from  $1 \times 10^{19}$  at the implant mask edge to say,  $5 \times 10^{18} \text{ cm}^{-3}$  at some distance  $\Delta w_R$ , the resistance increases ten-fold. The overall resistance is thus determined by the region of high doping. A  $1 \times 10^{19} \text{ cm}^{-3}$  is much larger than the doping in the drift collector. The capacitive straggle should be defined as the

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point when the doping falls to 10% of the collector doping or the point at which lateral Si concentration falls below the active Fe doping, whichever is lower.

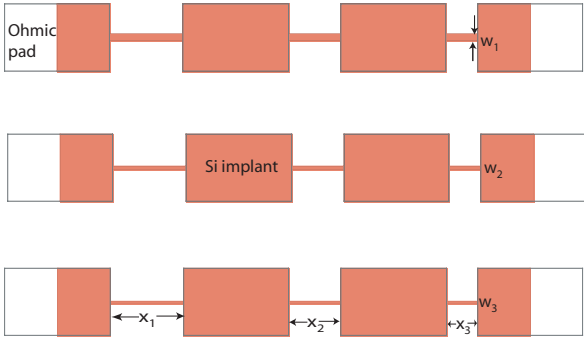


Figure 5.21: Straggle TLMs with varying spacing  $x$ , and width  $w$ – the shaded regions are implanted with Si, and ohmic contacts are formed over these

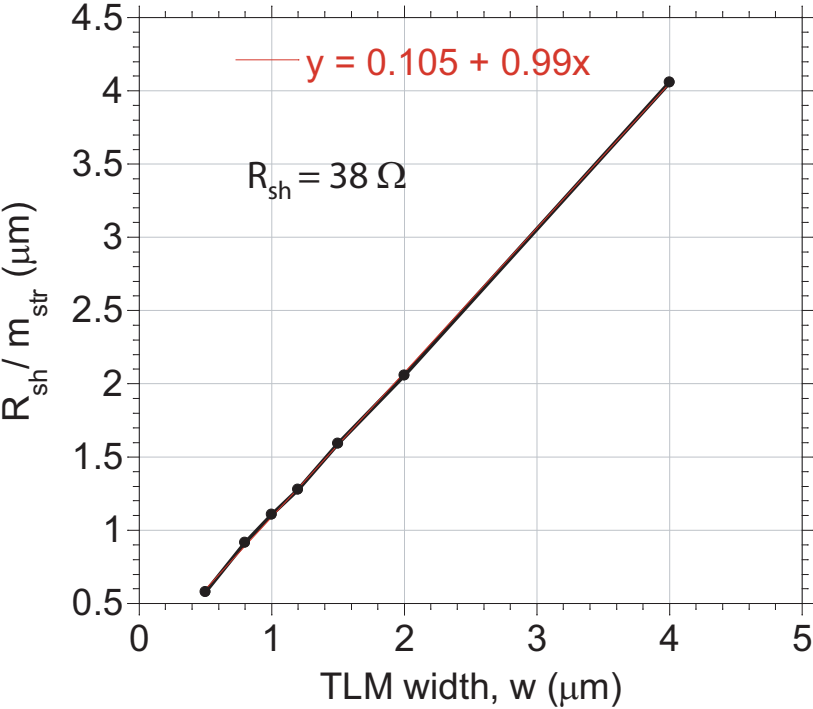


Figure 5.22: Resistive straggle of Si pedestal implant in Fe activated layer



Table 5.4: Contact and sheet resistivities

	Sheet resistance $\Omega/\square$	Contact resistivity $\Omega \cdot \mu\text{m}^2$	Horizontal resistance $\Omega \cdot \mu\text{m}$
Emitter	28	$< 1$	3
Base	1300	$< 3$	$< 50$
Collector	24-26	3000	350
Pd/Ti/Pd/Au contacts for all layers			

### 5.3 Large area device results

The RBS data in Fig. 5.18 is encouraging proof that good quality crystalline growth is feasible on this triple implanted substrate. Similar to the growth for implanted subcollector DHBTs, the drift collector, base and emitter are grown with the same layer structure as in Table 4.5. There is no pedestal layer, therefore this large area device is representative of the transistor in the extrinsic region under the base ohmic. The collector contact is formed by etching down  $\sim 0.3 \mu\text{m}$  to place the contact pads on  $N^{++}$  Si implanted subcollector. The sheet resistance of the collector, base and emitter of these devices in Table 5.4 are very similar to fully epitaxial layers grown at UCSB. Also excellent growth morphology is observed (not shown). The large area DC results are shown in Fig. 5.23. These indicate that there is no gain in this transistor due to current blocking by the SI region below the drift collector. Only at very high collector base biases, does current start to flow as also evident from the diode leakage in Fig. 5.24. The Gummels in Fig. 5.23a show that there is no current gain when  $V_{cb} = 0\text{V}$  and  $\beta \sim 1$  when  $V_{cb} = 1\text{V}$ . The CV measurements in Fig. 5.25 indicate that the capacitance at zero bias is  $0.5 \text{ fF}/\mu\text{m}^2$ . The capacitance of a fully epitaxial HBT with identical layer structure but epitaxially grown 300 nm  $N^{++}$  subcollector is also shown and is  $\sim 1.1 \text{ fF}/\mu\text{m}^2$ . Thus there is a 55% decrease in capacitance due to the SI region at zero bias. This corresponds to a depletion thickness of 220 nm, or a thickness of the SI region of  $\sim 120$  nm. The capacitance measurements beyond 0.5 V for the DHBT grown on the Fe and Si implanted substrate were clouded by high parallel conductance and is not shown. The offset voltage is higher since higher  $V_{cb}$  is required to remove the current blocking due to Fe. The high knee voltage is due to the large resistance in the SI Fe implanted region. From Fig. 5.26, the  $V_{cb}$  is over 1V only when  $V_{cb}$  is over 2V. A large  $V_{cb}$  is needed for the base collector junction of this device to be not forward biased. This explains the large offset and knee voltages

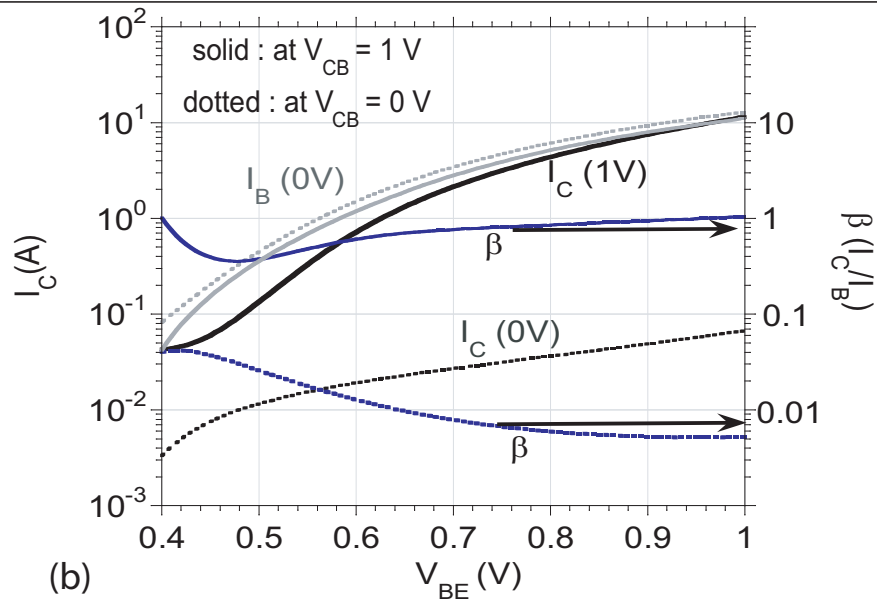
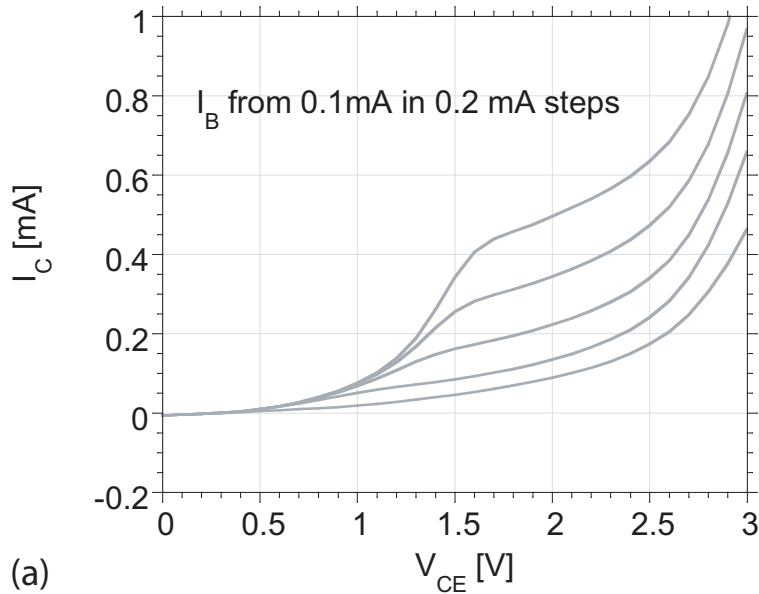


Figure 5.23: DC measurements of large area devices on Si and Fe implanted substrate – representative of extrinsic DHBT

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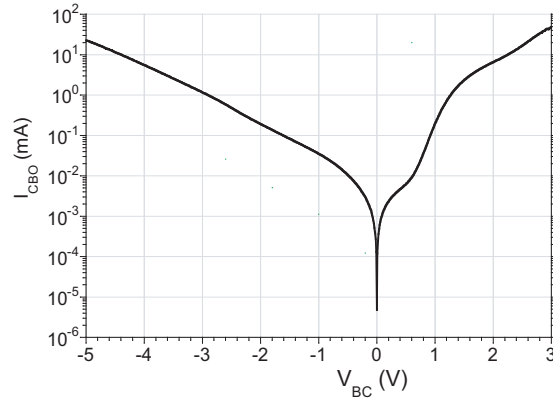


Figure 5.24:  $I_{CBO}$  of large area devices on Si and Fe implanted substrate – representative of extrinsic DHBT

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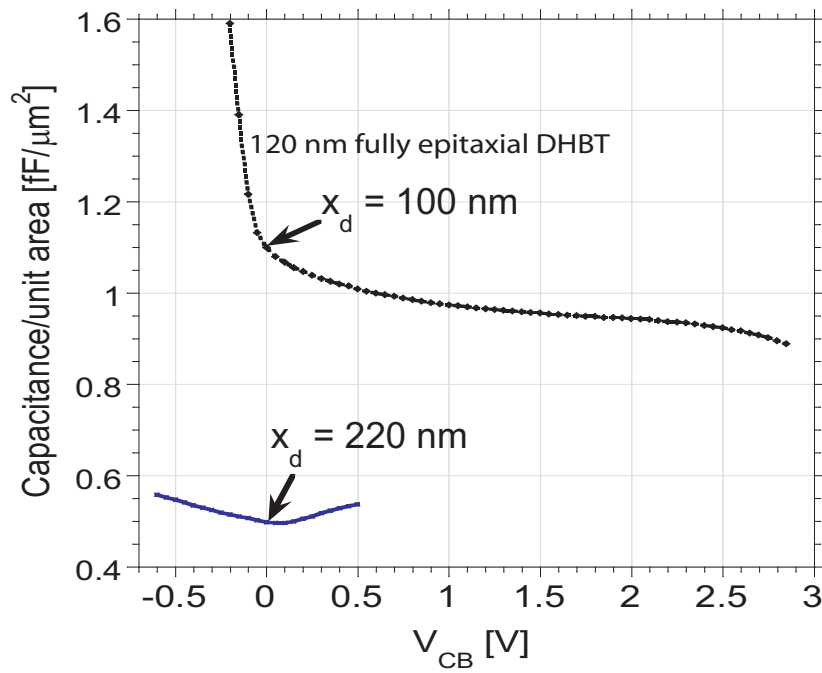


Figure 5.25: C-V measurements of large area devices on Si and Fe implanted substrate, showing a 55 % decrease in extrinsic capacitance compared to a standard epitaxial DHBT

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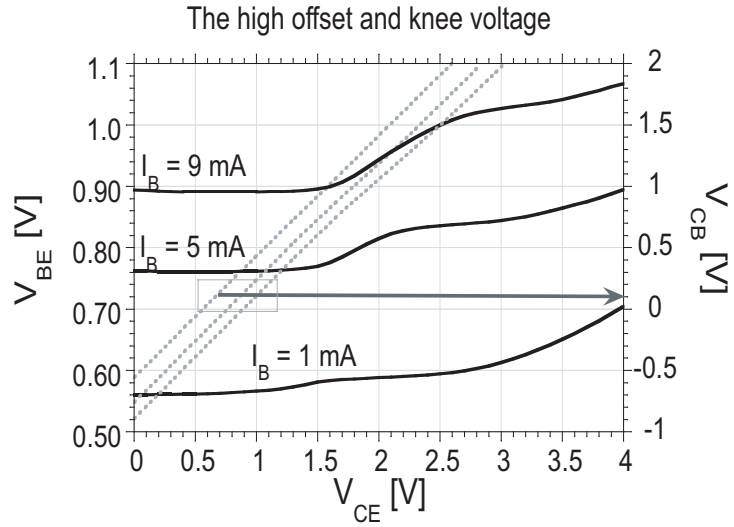


Figure 5.26:  $V_{be}$  and  $V_{cb}$  in the output characteristics

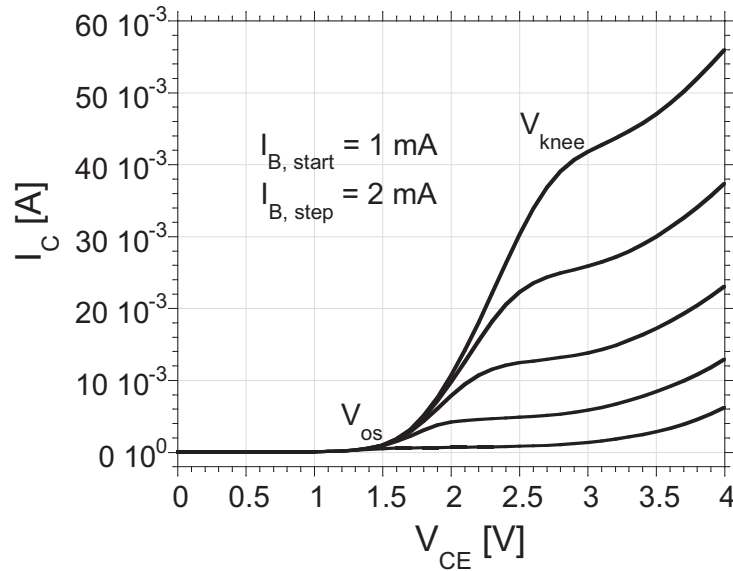


Figure 5.27: High  $V_{os}$  and high  $V_{knee}$  in the output characteristics

## 5.4 Device results - Implanted pedestal-subcollector DHBT

Large area devices have shown low leakage and excellent DC characteristics. Small area devices were processed with the identical layer structure shown in Table 4.5. The process has already been described in §4.4.1 and §5.2.

The TLM measurements of the collector and, of the pinched and non-pinched base TLMs are given in Fig. 5.29. The collector TLM structures is shown in Fig. 5.28. Thus the contact resistance measured is due to the ohmic contact and the resistance in the pedestal. This resistance from the TLMs is measured to be very high  $\approx 160\text{--}210 \Omega \cdot \mu\text{m}^2$ . This is partially due to the poor collector ohmic contact. The sheet resistance of the implanted  $N^{++}$  subcollector is  $\sim 21\text{--}22 \Omega/\square$ . The base sheet resistance is  $\sim 600 \Omega/\square$ , and the base contact resistivity is  $\sim 25 \Omega \cdot \mu\text{m}^2$ . The Isolation between ohmic pads separated by  $10 \mu\text{m}$  is xx pA/ $\mu\text{m}$  as seen in Fig. ?? For the

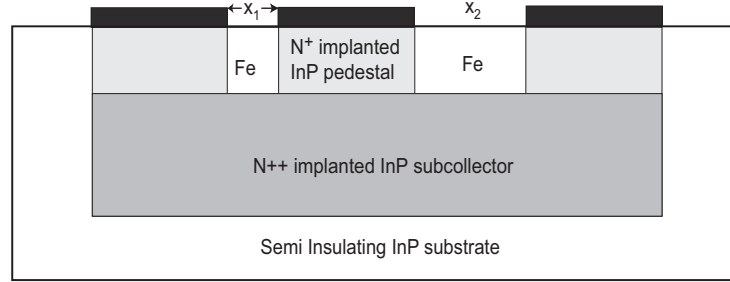


Figure 5.28: Collector TLM structure for the implanted pedestal subcollector DHBT

device with  $A_{jBE} = 0.65 \times 4.3 \mu\text{m}^2$ ,  $A_E = 0.7 \times 5.25 \mu\text{m}^2$  and active base collector area  $1.3 \times 5.25 \mu\text{m}^2$ , Fig. 5.30 is a plot of the DC  $I_C - V_{CE}$  characteristics. The maximum current gain at  $V_{cb} = 0$  is  $\sim 35$ . The Gummel curves are plotted for  $V_{cb} = 0\text{V}$  and  $V_{cb} = 0.3\text{V}$  Fig. 5.31. The collector and base ideality factors are  $\eta_c = 1.26$  and  $\eta_b = 1.77$ . The leakage current  $I_{CEO}$  from the gummel curves is  $\approx 1 \text{ nA}$  at  $V_{cb} = 0.3\text{V}$ . The DC current gain of the transistor  $= I_c/I_b$  from the gummel curves is shown in Fig. ?. The HBT's gain compression point is  $\approx 4 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0\text{V}$  and  $5 \text{ mA}/\mu\text{m}^2$  at  $V_{cb} = 0.3\text{V}$ . At high current densities,  $V_{CB,i} = V_{CB, applied} - I_C R_c$ . From the TLMs and the geometry of the transistor,  $R_c$  for this transistor should be  $\approx 11 \Omega$ . At  $I_c \approx 15\text{mA}$ ,  $V_{CB,i} = V_{CB, applied} - 0.3\text{V}$ . The base collector junction is forward biased till  $V_{CB, applied}$  is  $\approx 0.3\text{V}$ . The collector resistance is extracted from the  $I_C - V_{CE}$  characteristics of the transistor at saturation, using the Fixed  $\beta$  method

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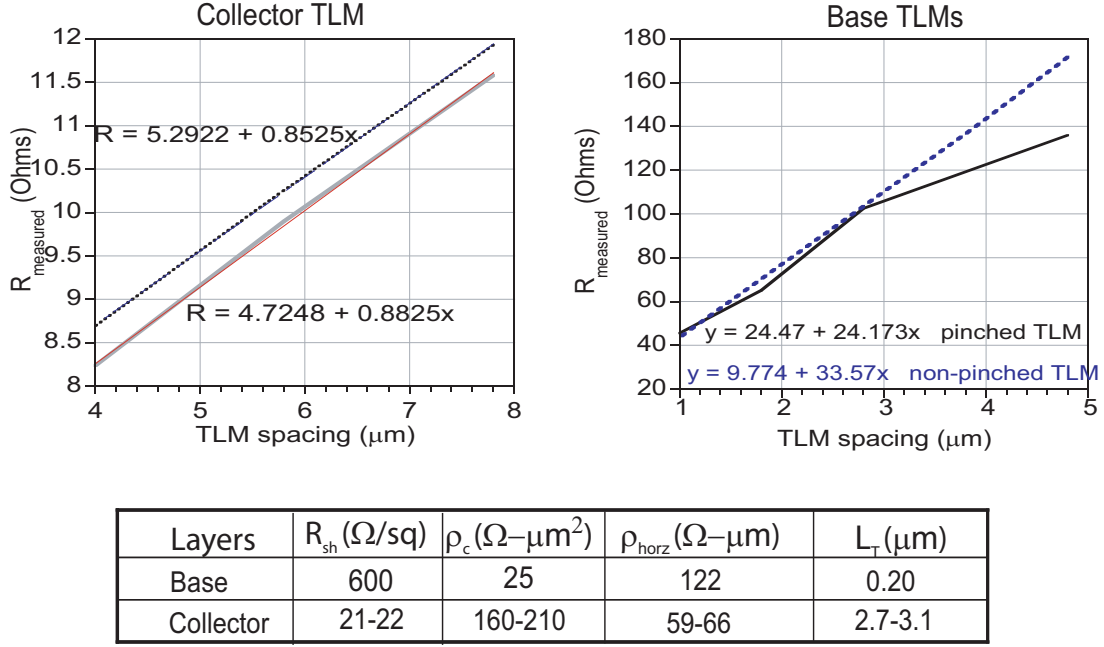


Figure 5.29: Base and Collector TLM measurements

and is plotted in Fig. ???. The emitter resistance,  $R_{ex} \approx 4 \Omega$  (from Fig. 5.35, the collector resistance is determined to be  $\approx 32-34 \Omega$ , consistent with the value calculated from TLM measurements. This is very high compared to the value of  $1-3\Omega$  that is typically obtained and is partly due to the poor ohmic contact to  $N^{++}$  InP.

The resistance as seen in the emitter,  $R_{ex} + R_{bb}/\beta$  is derived from gummels and plotted as function of current density Fig. 5.35. From TLM's measured,  $R_{bb}/\beta \approx 1 \Omega$  and therefore  $R_{ex} \approx 4 \Omega$ . Also plotted in Fig. ?? is the thermal resistance,  $\Theta_{th} \cdot \phi$  (also referred to as  $R_{th}$ ). The thermal-electric feedback coefficient  $\phi$  from [8], is  $0.85 \text{ mV}/^\circ \text{K}$  at  $5 \text{ mA}/\mu m^2$ . The thermal resistance at  $5 \text{ mA}/\mu m^2$  is therefore  $2.7 \text{ }^\circ \text{K}/\text{mW}$ . Shown in Fig. 5.36 are  $I_{CEO}$  and  $I_{CBO}$  of a triple mesa DHBT and an implanted pedestal-subcollector DHBT. The breakdown voltages are defined at  $I_c = 1 \text{ mA}$ .  $BV_{CEO} = 6.8\text{V}$  and  $BV_{CBO} = 7.6\text{V}$ . For a fully epitaxial, standard mesa DHBT, the corresponding voltages are  $BV_{CEO} = 5.5\text{V}$  and  $BV_{CBO} = 6.8\text{V}$  at  $I_c = 1\text{mA}$ . The breakdown enhancement in these implanted pedestal-subcollector devices is due to lower surface breakdown due to suppression of surface states. The safe operating area is a more relevant than breakdown voltage for digital logic circuits and is seen in Fig. 5.37. The maximum operating power density is seen to be  $\sim 15 \text{ mW}/\mu m^2$ .

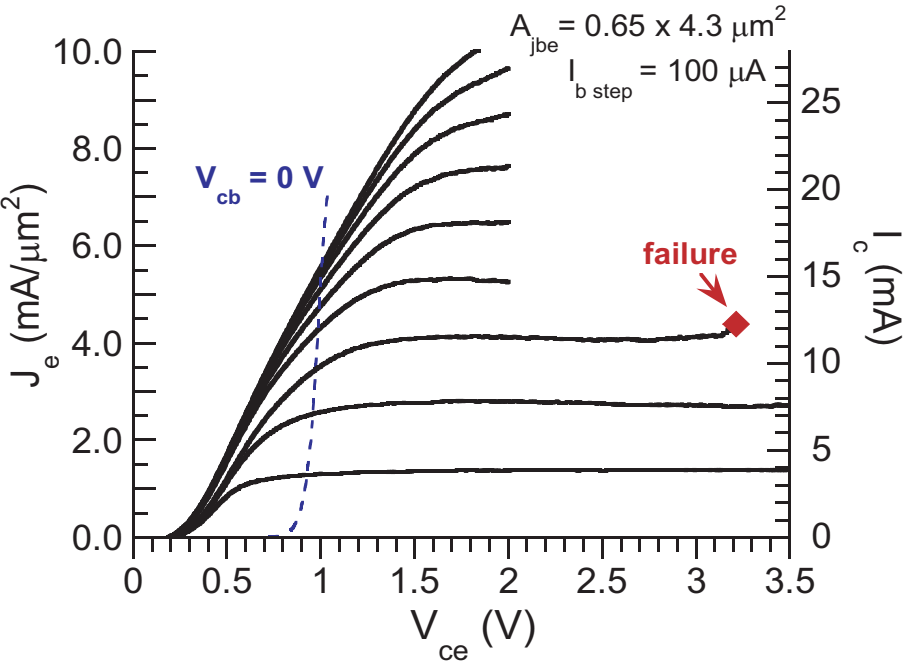


Figure 5.30: DC  $I_C - V_{CE}$  characteristics

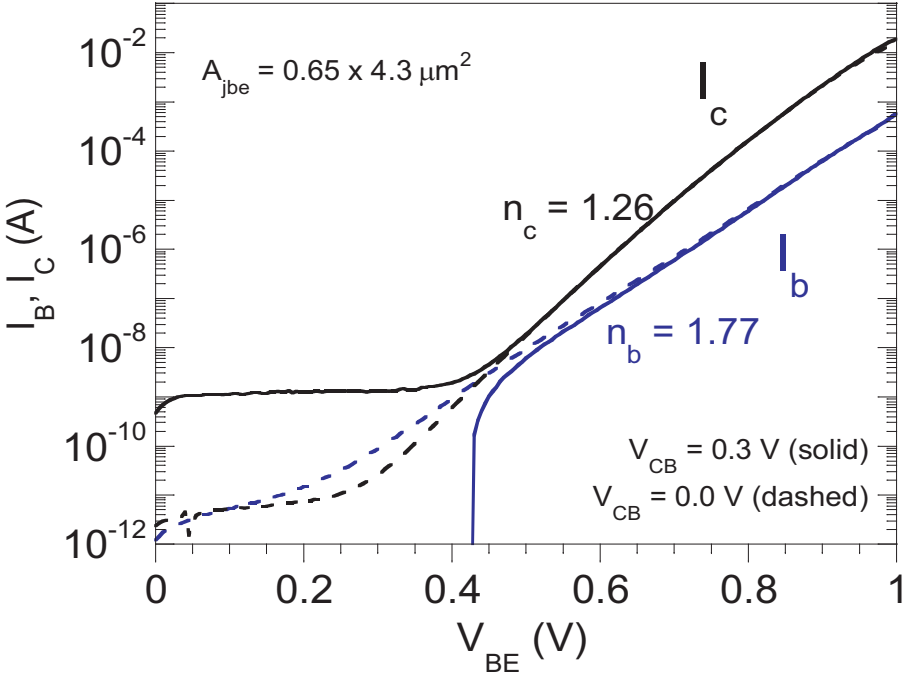


Figure 5.31: Gummel curves at  $V_{cb} = 0\text{V}$  and  $0.3\text{V}$



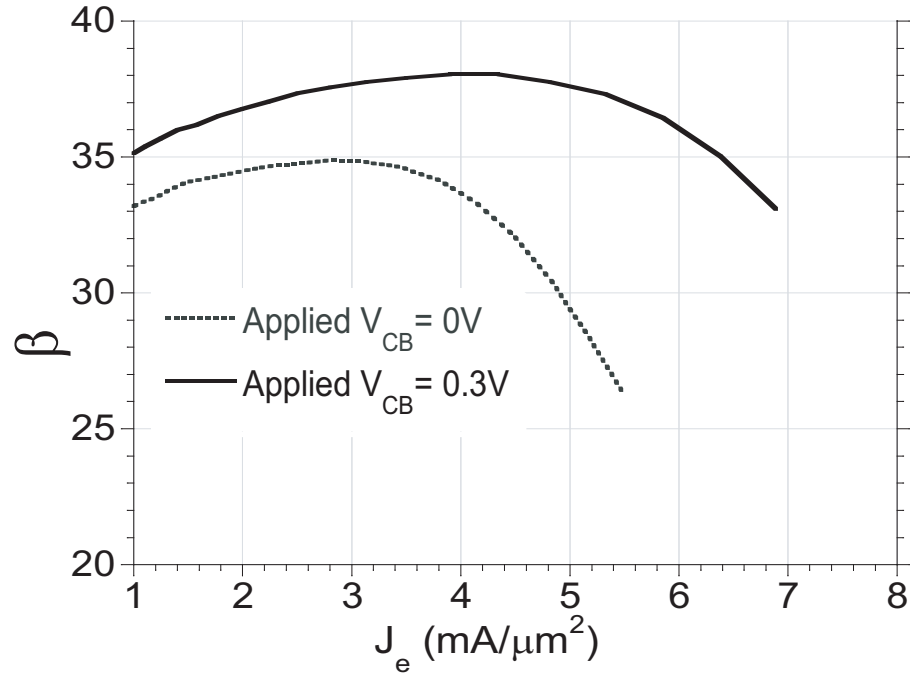


Figure 5.32: DC gain of the HBT at  $V_{cb} = 0\text{V}$  and  $0.3\text{V}$

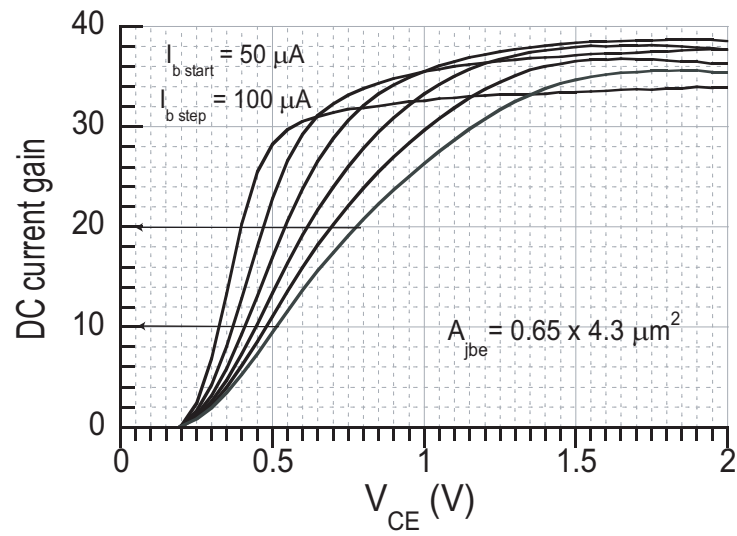


Figure 5.33: Extraction of Collector resistance from saturation characteristics

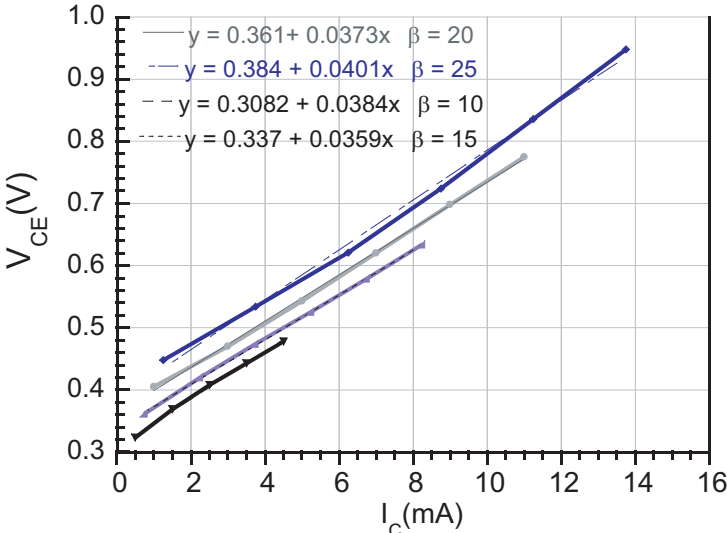


Figure 5.34: Collector resistance at various  $\beta$ 's

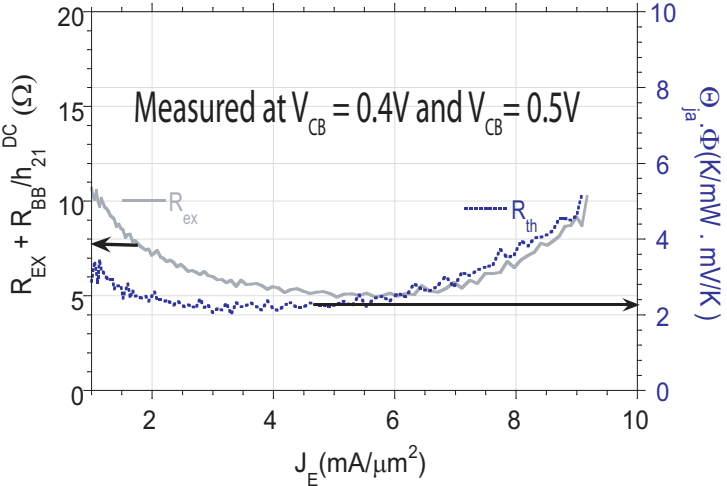


Figure 5.35: Extraction of Emitter resistance, and thermal resistance

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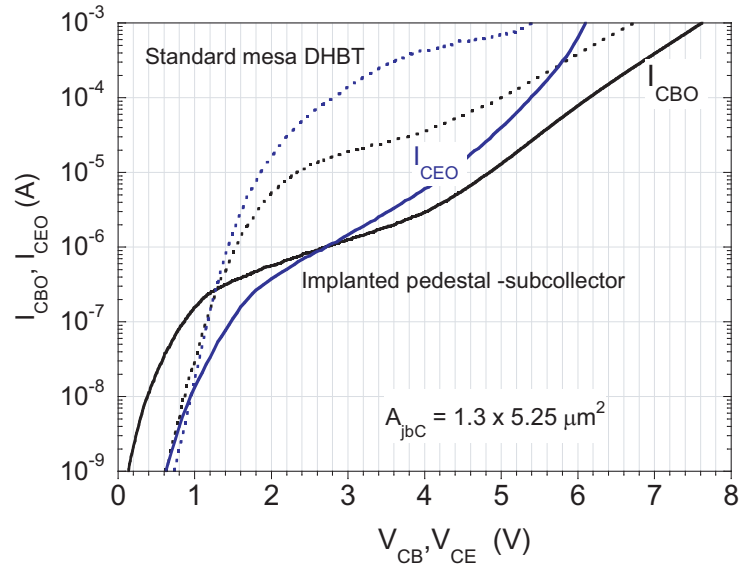


Figure 5.36: Leakage currents in the device

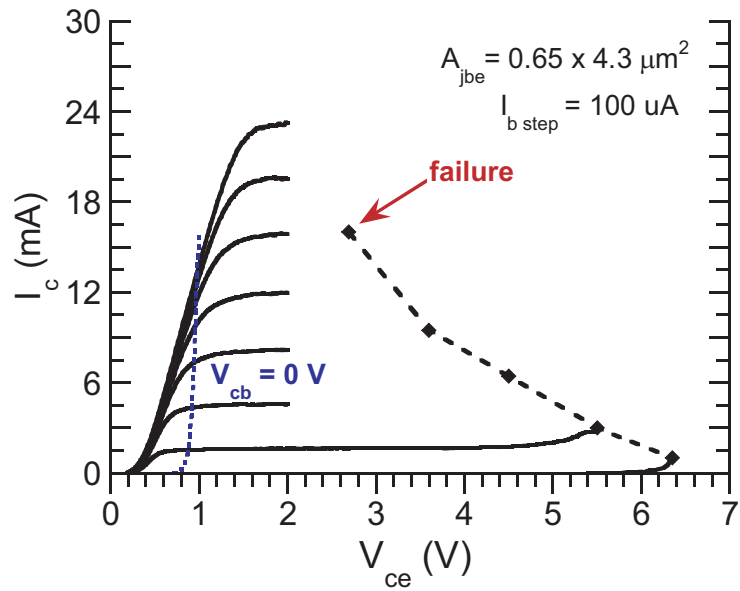


Figure 5.37: Safe Operating area of the device

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DC-45 GHz S-parameter measurements are carried out after performing an off wafer Line-Reflect-Reflect-Match calibration on an Agilent 8510C network analyzer. On-wafer open and short circuit pad structures identical to the ones used by the devices are measured after calibration in order to de-embed their associated parasitics from the device measurements.  $f_{max}$  is determined from extrapolation through a least-square-fit between the transfer function,

$$U(f) = \frac{U_{DC}}{1 + U_{DC} \cdot (f/f_{max})^2} \quad (5.4.0)$$

to the measured microwave gain U at measured frequencies.  $1/f_\tau$  is the slope of  $\text{Im}(1/h_{21})$  as explained in Chapter 4. A maximum 352 GHz  $f_\tau$  and 403 GHz  $f_{max}$

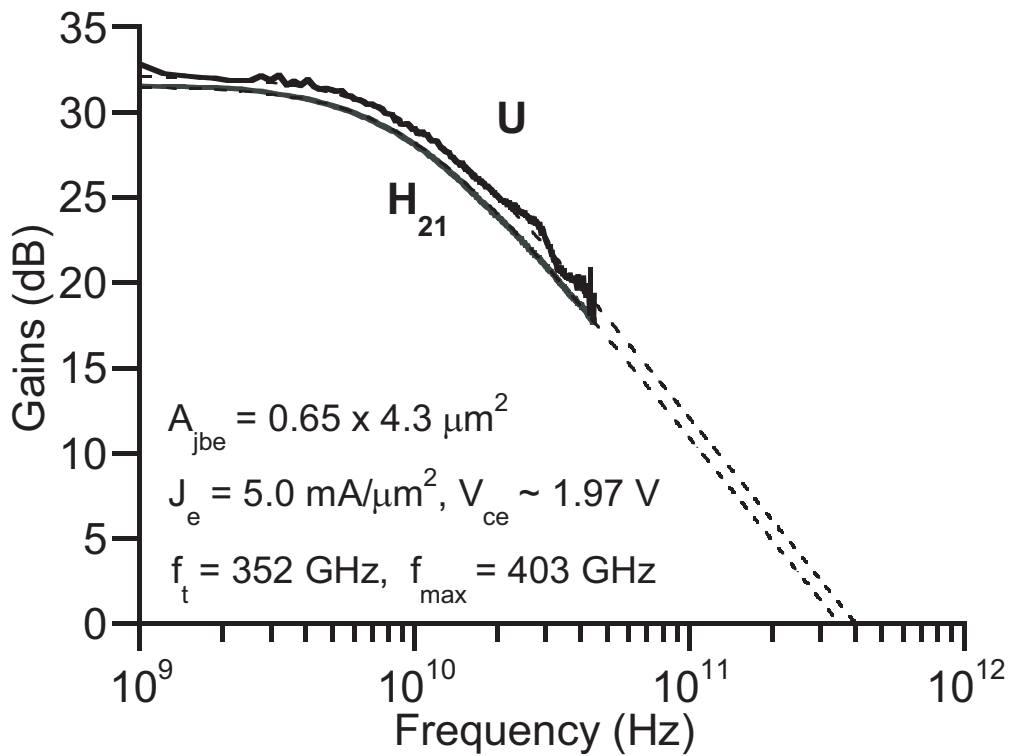


Figure 5.38: Microwave gains of Si implanted subcollector DHBTs

is demonstrated at  $I_c = 14 \text{ mA}$  and  $V_{cb} = 1.96 \text{ V}$  ( $J_e = 5 \text{ mA}/\mu\text{m}^2$ ,  $C_{cb}/I_c = 0.43 \text{ ps/V}$ ). These are lower than the standard, fully epitaxial, triple mesa DHBT. The collector resistance is much higher ( $32 \Omega$  vs.  $2\Omega$ ).

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The capacitance voltage characteristics of the implanted subcollector DHBTs with Fe at  $I_c = 0$  mA, are plotted and compared with the standard mesa equivalent in Fig. 5.39. From the figure, The implanted subcollector DHBTs with Fe exhibit a  $C_{cb}$  reduction of  $\approx 3.2$  fF over the entire measured range of bias voltages due to elimination of the charge dipole at the growth interface. At full depletion, this corresponds to a 25-27% decrease in  $C_{cb}$ . This is lower than expected reduction in  $C_{cb}$  and is attributed to lateral implant straggle. Here,  $A_{jbc} = 1.3 \times 5.25 \mu\text{m}^2$   $A_{pad} = 2.5 \mu\text{m}^2$ . The pedestal stripe as defined on the photo mask is  $0.9 \times 5.25 \mu\text{m}^2$ . The length of the subcollector as defined on the photo mask is  $5.25 \mu\text{m}$ . The voltage drop  $I_c \times R_c$  is significant and the collector potential must be progressively increased as  $I_c$ , to fully deplete the drift collector. The lower  $f_\tau$  and  $f_{max}$  demonstrated by the implanted HBTs compared to their triple-mesa counterpart is due mostly to the increased delay associated with the terms  $kT/qI_c (C_{cb} + C_{je})$  and  $R_c C_{cb}$ . Fig. 5.41 shows  $C_{cb}/I_c$  as a function of bias.  $C_{cb}/I_c$  is an important metric for digital logic circuits.

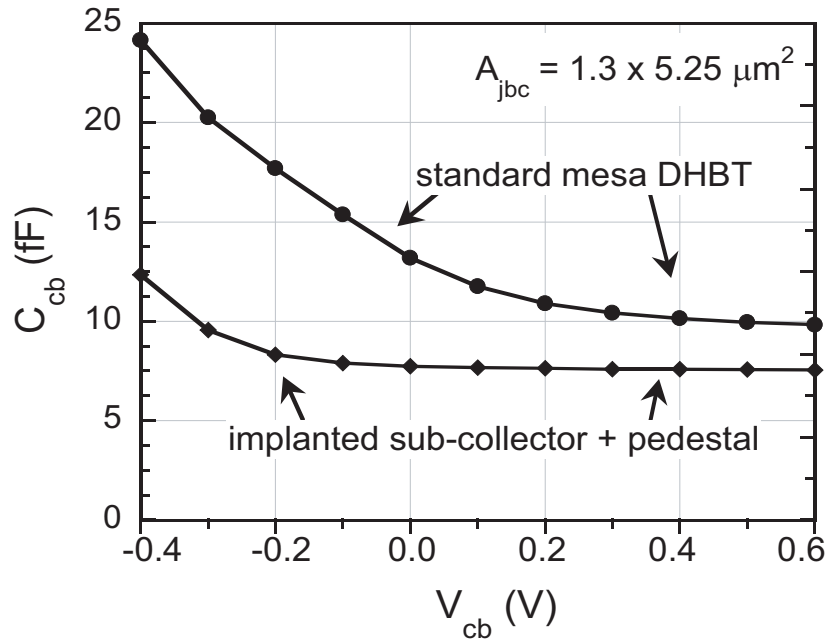


Figure 5.39: Variation of  $C_{cb}$  with  $C_{cb}$  at  $I_c = 0$  mA for a mesa HBT and the pedestal HBT

The slightly lower  $C_{cb}$  reduction than expected, is due to lateral and longitudinal implant straggle. The DC results compare with the standard mesa DHBT and hence

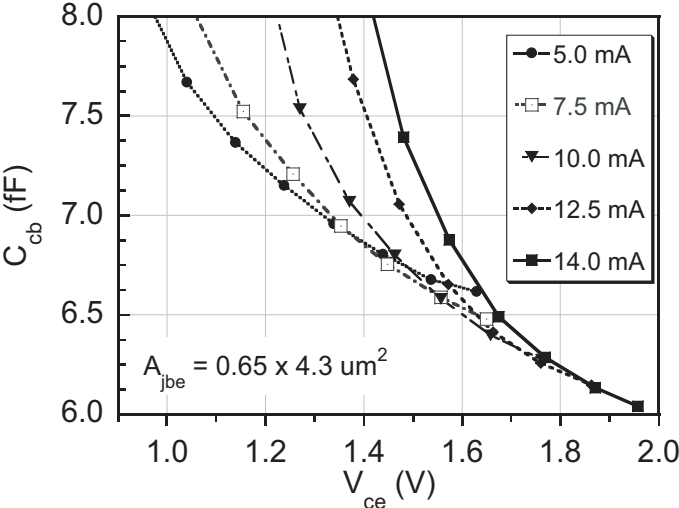


Figure 5.40: Variation of  $C_{cb}$  with  $V_{cb}$  at different  $I_c$  to show the effect of high series collector resistance

indicate the feasibility of manufacturing microwave HBTs on the triple implanted substrates. The high collector contact resistance has resulted in lower  $f_\tau$  and  $f_{max}$  compared to [9].

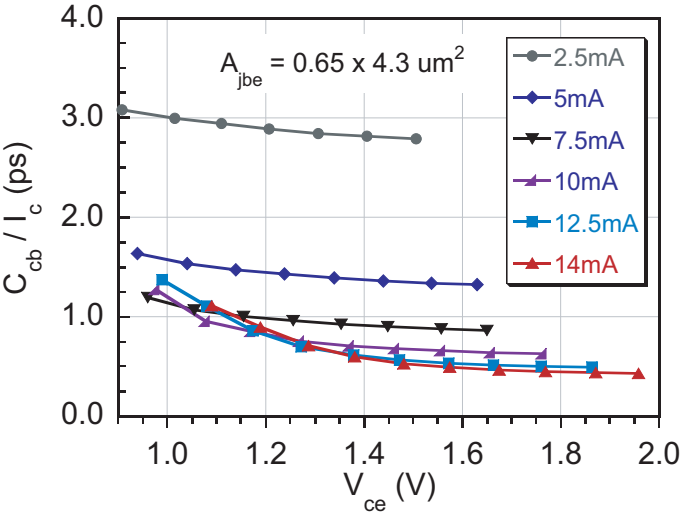


Figure 5.41:  $C_{cb}/I_c$  as a function of bias

## 5.5 Discussion

These first generation fully implanted pedestal-subcollector DHBTs have record bandwidths for implanted collector DHBTs. However, they suffer from some problems

- Large collector resistance
- less than expected  $C_{cb}$  reduction

These issues will be discussed in the following sections and solutions suggested

### 5.5.1 Pedestal vertical height and lateral straggle

In order to evaluate the height of the pedestal, C-V measurements are carried out on devices with no  $N^{++}$  pedestal implant. These have the  $N^-$  collector directly atop the SI Fe implanted region and therefore resembles the extrinsic portion of the device,  $\hat{C}_{cb} = 0.54 \text{ fF}/\mu\text{m}^2$ . This is compared with a device, where the  $N^{++}$  pedestal and subcollector extends throughout the device so that it resembles a mesa structure.  $\hat{C}_{cb}$  for this mesa-like device is  $1.1 \text{ fF}/\mu\text{m}^2$  and the  $\tau_c = 120 \text{ nm}$ . Therefore the height of the pedestal is deduced to be  $\sim 120 \text{ nm}$ .

At full depletion,  $C_{cb}$  is measured as for various devices with different pedestal geometries and plotted as function of pedestal width to determine the lateral pedestal straggle.

$$\frac{C_{cb}}{\epsilon/T_c} \cdot \frac{1}{L_e} = 0.5 \cdot (W_{BC} + W_p) + \Delta W_p \quad (5.5.0)$$

where  $\Delta W_p$  is the capacitive lateral straggle on each side of the pedestal. This normalized capacitance is plotted vs.  $W_p$  and is shown in Fig. 5.43. From this, the straggle on either side is  $\sim 0.2 \mu\text{m}$  and is responsible for the less than anticipated reduction in  $C_{cb}$ . This calculation ignores the longitudinal straggle due to the subcollector implant.

The simplest way to address the stragging problem is to reduce the dimension of the mask. However, less than  $0.5 \mu\text{m}$  features are hard to define using the optical stepper system at UCSB. Therefore one method of reducing the  $Si_xN_y$  implant mask dimensions is shown in Fig. 5.44. However, the resistive straggle is seen to be  $\sim 0.05 \mu\text{m}$ . Therefore reducing the mask dimension laterally, reduces the capacitance but adds to the vertical pedestal resistance. SIMS data on a test wafer, shows no diffusion of the Fe after growth.



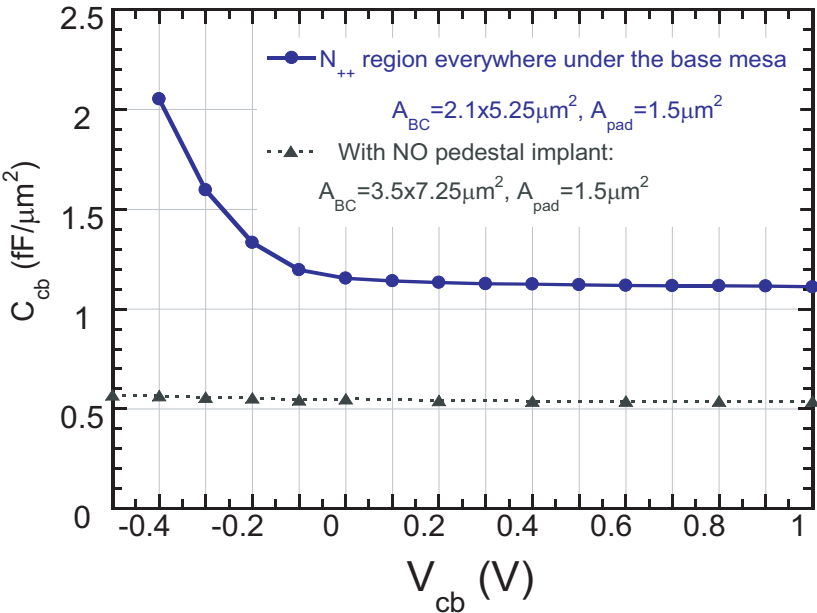


Figure 5.42: C-V measurements of a device without an  $N^{++}$  pedestal and one with  $N^{++}$  subcollector and pedestal everywhere in the device footprint so that it resembles a mesa geometry.

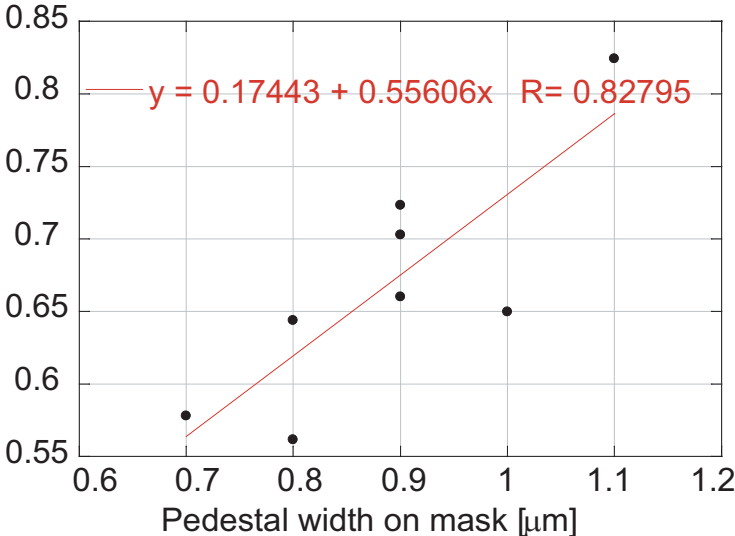


Figure 5.43: Normalized C vs. pedestal width to determine capacitive straggle due to implant

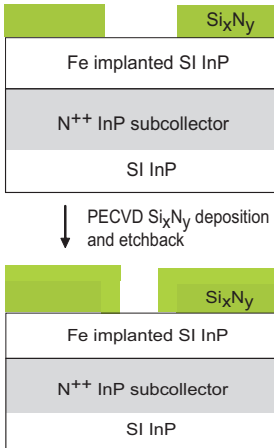


Figure 5.44: To obtain submicron features for implant mask...

CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

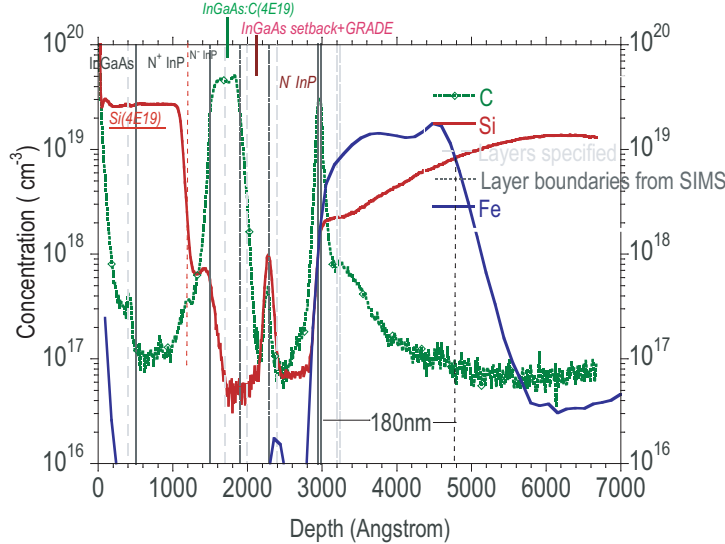


Figure 5.45: SIMS of the HBT layer stack after growth on implanted layers. There is no pedestal implant here

### 5.5.2 Excess Collector Resistance

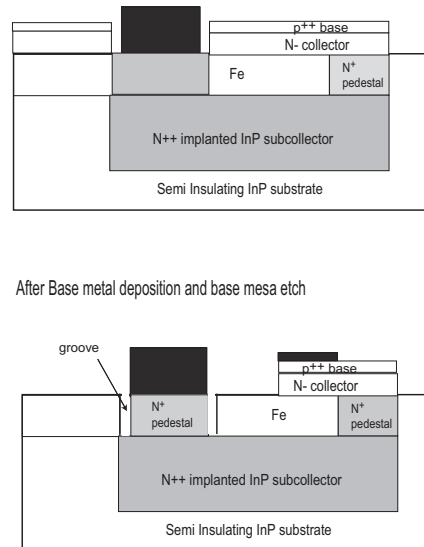


Figure 5.46: Reverse collector ohmic process

As with the implanted subcollector DHBTs, these DHBTs too suffer from poor collector contact resistance due to insufficient anneal. A new process is proposed, where the collector metal is deposited by recess etching through the base and collector layers. The collector metal does not perfectly fill this recess but instead resembles Fig. 5.46a. When the base mesa is etched, the semiconductor next to the collector contact is etched as well, as seen in Fig. 5.46b. However, this does not in any way impact the implanted pedestal-subcollector DHBT as the semiconductor etched away is Fe implanted InP layer. Therefore the reverse collector ohmic process is feasible for implanted pedestal subcollector DHBTs even with the current photo masks by merely reversing the deposition of the base and collector ohmics. This now permits a much higher anneal for the collector ohmic, reducing the contact resistivity to  $20\text{-}25 \Omega \cdot \mu\text{m}^2$ .

From §5.4, it is seen that the calculated  $R_c$  from TLMs is  $11 \Omega$ . This does not explain the very high resistance seen in the devices of  $\sim 32 \Omega$ . Several devices were measured and the excess  $R_c$  values tabulated in Fig. 5.47.

These suggest that the excess resistance is due to the pedestal itself. Special test structures were placed on another wafer implanted similarly. The pedestal TLM is a special test structure (Fig. 5.48), where only the Si pedestal implant is carried out in a background of Fe already implanted and activated. From these, the sheet

CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

$W_p(\mu\text{m})$	$L_p(\mu\text{m})$	$R_c(\Omega)$	$R_c$ from TLM( $\Omega$ )	Excess $R_c$ ( $\Omega\text{-}\mu\text{m}$ )
0.8	5.25	34.4	12	117.6
1	7.25	23.2	8.6	105.9
Mesa	Mesa	29.1	11.6	91.9
0.9	4.25	35.3	14.7	87.5
1.3	5.25	35.2	11.6	128.6

Figure 5.47: Excess Collector resistance of devices of various geometry

resistance of the pedestal is deduced to be  $\sim 38 \Omega/\square$ , exactly as measured earlier. The vertical contact resistance is now entirely due to the ohmic contact and is seen to be  $\sim 25 \Omega \cdot \mu\text{m}^2$ . The pedestal-subcollector TLM in Fig. 5.48 measures a contact resistivity of  $60 \Omega \cdot \mu\text{m}^2$ . Since the pedestal resistivity is low, this means that there is a resistance of  $35 \Omega \cdot \mu\text{m}^2$  at the interface between the pedestal and the subcollector.

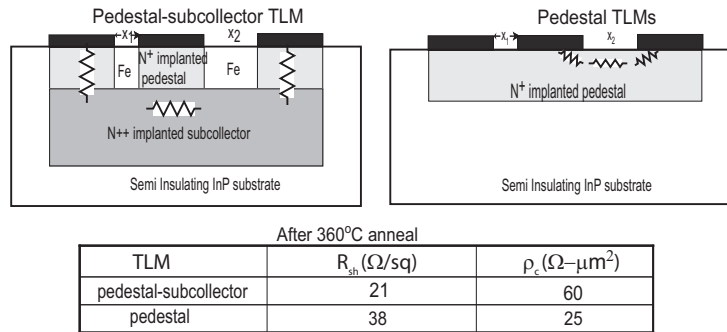


Figure 5.48: Special test structures to find the source of excess  $R_c$

This is attributed to the large Fe pile up seen in §5.2.1. The  $R_c$  is not due to the narrow pedestal dimensions but instead due to insufficient Si at the region of Fe pileup. This is evident from the common emitter IVs of two devices in Fig. 5.49, one with a narrow pedestal and one where the pedestal and subcollector are present over the entire device footprint.

As discussed in Chapter 3, the Fe preferentially piles up in the region of most damage. RBS does not detect defects below  $\sim 1 \times 10^{20} \text{ cm}^{-3}$ . This is still a large number of defects. e implant is performed at a lower dose ( $1 \times 10^{14} \text{ ions/cm}^2$ ) and

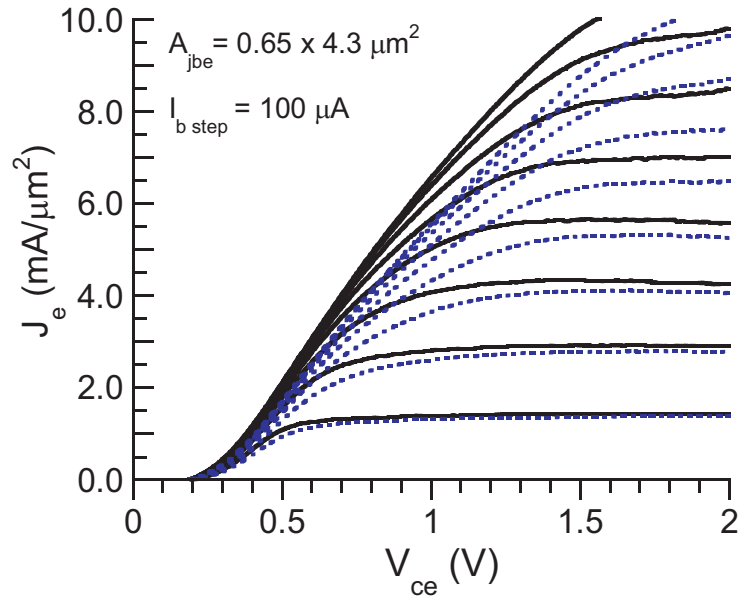


Figure 5.49: SIMS of the HBT layer stack after growth on implanted layers. There is no pedestal implant here

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lower energy (100 keV) to check if there is a dose dependence of such damage. There is no evidence of Fe pile up at these implant conditions. CV measurements on Schottky diodes are performed to check the thickness of the Fe compensated layer for these conditions. This is shown in Fig. 5.51

While these first generation HBTs suffered from certain problems, it is shown that these can be resolved.

CHAPTER 5. IMPLANTED PEDESTAL-SUBCOLLECTOR DHBT

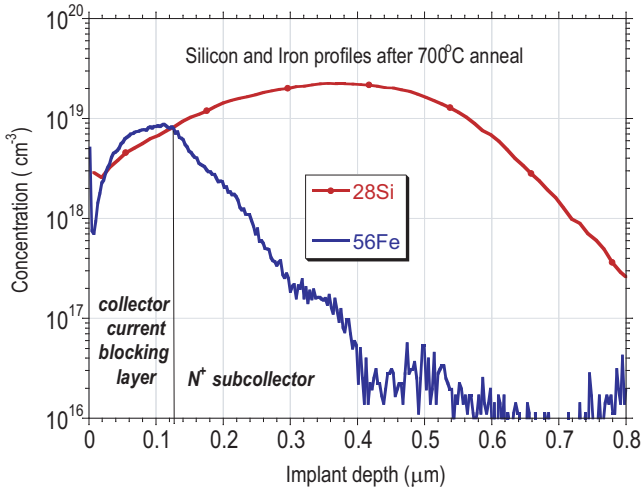


Figure 5.50: Distribution of Fe and Si after anneal

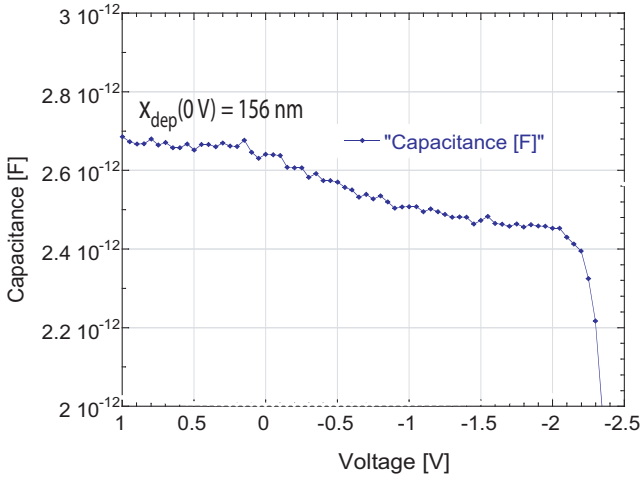


Figure 5.51: CV measurements of Schottky diodes to determine the depletion thickness for these implant conditions

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# 6

## Conclusions

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**I**MPLANTED subcollector and pedestal DHBTs were demonstrated for the first time in an all implanted, manufacturable process. Record bandwidths for DHBTs with implanted collectors have been demonstrated. These were incorporated in high speed CML and ECL logic circuits that have demonstrated over 20% improvement in digital logic speed over conventional mesa DHBTs.

### 6.1 Accomplishments

As the current mesa InP DHBT is reaching its scaling limit, new process modules were characterized and added to further the limits of such devices [1]. The parasitic collector base capacitance is especially an important hurdle for high bandwidth analog circuits and high speed digital circuits. A manufacturable technology that enable submicron collector scaling of InP HBTs utilizing ion implantation is proposed and developed. For this purpose, Si and Fe implants in InP are extensively studied and characterized. With the lithographic dimensions feasible today, the device design has been tailored for high speed digital circuit performance.

Modern mesa DHBTs have an extrinsic access pad that contributes significantly to the collector base capacitance. This capacitance is non-scalable and becomes important as device length is reduced. A selectively implanted Si subcollector is suggested in Chapter 4 to eliminate this base access pad capacitance. However, devices fabricated did not show any reduction in collector base capacitance at low bias voltages due to the presence of an N-type interface charge at the growth surface. A new method of eliminating this interface charge using Fe implants is proposed and demonstrated to work effectively over a wide range of charge densities [2]. Implanted subcollector DHBTs utilizing selective Si implants for isolation of the subcollector and Fe implants for eliminating the interface charge are developed. These devices is shown to almost completely eliminate this base access pad capacitance and have cut-off frequencies in excess of 350 GHz [3]. This technology enables for low power, high speed digital circuits. The bandwidth of the devices are limited by

## CHAPTER 6. CONCLUSIONS

the excessive base and collector resistance due to poor ohmic contacts. Improving the ohmic technology should yield superior, high speed devices in a simple process.

Further collector scaling is made possible through a novel fully implanted pedestal-subcollector technology described in Chapter 5 . An implanted, buried subcollector is formed in SI InP substrate by high energy Si implants. Fe is used to compensate the top  $0.2 \mu\text{m}$  of Si to make this region Semi-Insulating. An additional set of Si implants then form an  $N^{++}$  pedestal link to the subcollector. Such a topology requires no extra growths and is very planar. By tailoring the width of the pedestal, the collector can be scaled independent of the base mesa dimensions. Several experiments and measurements are carried out to completely characterize this device structure. The sheet resistance of the subcollector and pedestal are designed to be low, so as to not hurt the collector resistance. The implant and anneal conditions are developed to result in a crystalline semiconductor before growth. Standard mesa devices can then be fabricated after growth. Thus this process is very modular. Microwave devices fabricated, demonstrated  $f_{\tau}$ 's of over 350 GHz and  $f_{max}$  of over 400 GHz [4]. The breakdown voltages are higher than triple mesa DHBTs due to lower active surface states. The  $C_{cb}$  reduction was  $\sim 30 \%$ . The large collector resistance and lateral straggle due to the pedestal implant has limited the performance of this device. The collector resistance is higher than a standard mesa DHBT, partly due to a poor ohmic contact and partly due to Fe piling up at the interface between the pedestal and the subcollector, causing a vertical resistance at this interface. This is due to the large implant dose of Fe. This has been solved by using a lower fluence of Fe where no such pile-up is observed (§5.5 in Chapter 5 ). The lateral straggle can be solved by reducing the dimensions of the implant mask, as discussed in §5.5 in Chapter 5 . With these two modifications, cut off frequencies in excess of 500 GHz are expected, at the current scaling node. As devices are laterally and vertically scaled,  $C_{cb}$  contributes increasingly to delay of analog and digital circuits. Such a technology should enable  $C_{cb}$  reduction in an elegant, high yield process.

Digital latches were fabricated using a similar collector pedestal process at Rockwell Scientific. These devices had cut-off frequencies of  $\sim 350$  GHz. Low power CML dividers having  $\Delta V_{logic}$  is 250 mV and effective loading resistance of  $100 \Omega - I_{data} = 2.5$  mA, are demonstrated [5]. The divider operation is fully static, operating from  $f_{clock} = 4$  GHz to 61.2 GHz while dissipating 27.1 mW of power in the flip-flop from a single -2.30 V supply. The power-delay product of this circuit is 113.0 fJ/latch. This is a record low power-delay product for an InP DHBT-based static frequency divider. Without the implanted collector technology, dividers in a regular mesa DHBT process clocked at a maximum of 51 GHz while consuming 29.2 mW of power. Thus a 20% improvement in logic speed is achieved. Ultra high speed CML logic circuits have also been demonstrated. With the collector pedestal technology, the maximum  $f_{clock}$  is  $\sim 135$  GHz, while dividers in the triple mesa process

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clocked at a maximum of 110 GHz. Initial testing of ECL static frequency dividers show promising results, with self oscillation frequencies in excess of 95 GHz, a 10 % improvement over the fastest logic circuits demonstrated [6]. These results have been obtained with first generation DHBTs with collector pedestals with the process still at its infancy. Very high bandwidth power amplifiers at over 300 GHz and digital circuits clocking at 200 GHz are feasible even in the near future.

### 6.2 Future work

The first demonstration of these implanted collector yielded devices with record bandwidths. However, the full potential of this process is limited by

- Large straggle due to pedestal implant
- Large vertical resistance in the pedestal collector due to Fe pileup
- Poor ohmic contacts to  $N^{++}$  InP

As discussed in §5.5, the contact resistivity to the  $N^{++}$  InP collector can be reduced by forming the alloyed collector ohmics before the base metal deposition, allowing a very high temperature anneal. The Fe pile-up is shown to be decreased at lower implant doses of Fe, while the pedestal implant mask can be redesigned to overcome the lateral implant straggle. By carefully choosing the implant doses of Fe and Si pedestal implant, these problems is shown to be overcome. Simulations predict device bandwidth in excess of 500 GHz with these enhancements.

The triple implanted pedestal subcollector process is very modular and can be inserted into many HBT technologies. Regrown HBTs have been demonstrated with  $0.3\mu\text{m}$  emitter features [7]. As seen in Chapter 2 , the emitter resistance also needs to be reduced with scaling generations. Utilizing very thin collectors and base layers, the implanted collector processes can be combined with emitter regrowth, to decrease the transit times and RC charging time constants. Initial simulations predict devices with  $f_{max}$  close to 800 GHz (Chapter 2 ). The emitter resistance is limited by the contact resistivity of the ohmic contact. *In situ* MBE deposited ErAs is known to form a perfect ohmic to InGaAs [8]. The implanted collector process can again inserted into a process utilizing ErAs ohmics for the emitter. All three processes can also be combined to yield HBTs with cut-off frequencies in excess of a THz and digital logic circuits clocking at over 300 GHz.

As seen in Chapter 5 , the Fe compensation of Si implant results in effective blocking of current till  $\sim 2\text{V}$ . This suggests a restructuring of the HBT, to form a collector-up HBT such that Fe implant is used for emitter scaling, as seen in Fig. ??.

## CHAPTER 6. CONCLUSIONS

A  $N^-/N^{++}$  template is grown and Fe is selectively implanted through the  $N^-$  emitter. Taking advantage of the lateral straggle of this implant, narrow emitter base junctions can be defined. The base and collector are then grown, and the device is formed in a similar triple mesa process as in [6]. The collector base junction is now very narrow and there is zero extrinsic capacitance. The emitter resistance is now a lateral resistance. The  $N^{++}$  emitter can be made thick to reduce the lateral access resistance in the semiconductor. This hurts device planarity, however, trench isolation techniques can be developed as in Fig. 6.2. The emitter resistance is, to first order, independent of the emitter base junction area.

$$C_{cb} \propto \frac{L_e W_c}{T_c} \quad (6.2.1)$$

$$R_{ex} \propto \frac{\rho_c}{L_e} \quad (6.2.2)$$

Hence, devices can be laterally scaled without significantly affecting the emitter resistance. This is particularly useful for high speed digital logic circuits.

With such enhancements described above, InP HBTs are poised to break the 1 THz barrier. These ultra high speed devices may find applications in high bandwidth communication links and sensing operations. The majority of the applications will remain unforeseen. Indeed, *the principal applications of any sufficiently new and innovative technology have always been applications that were created by that technology* [9].

CHAPTER 6. CONCLUSIONS

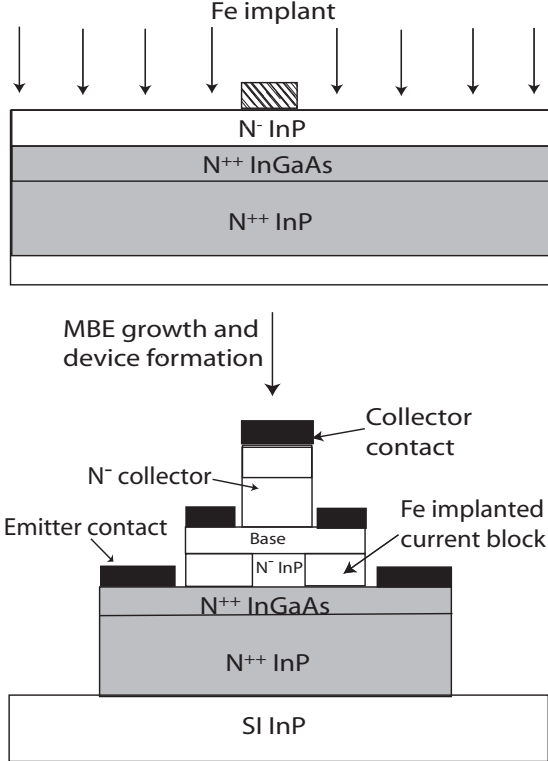


Figure 6.1: Proposal of a collector-up process with implants

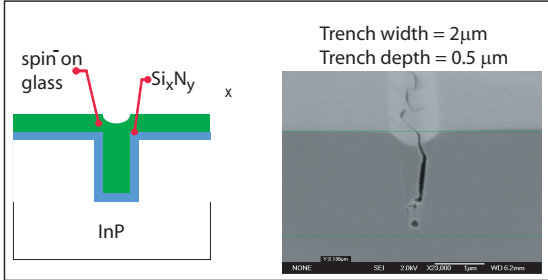


Figure 6.2: Trench isolation for InP HBTs

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## Implantation Analysis

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SEVERAL techniques exist to evaluate the implants. Various parameters relating to the implant need to be determined.

- Spatial distribution of the implanted ions to determine exact ion range and ion diffusion
- Crystallinity or defect density in the material due to the implant
- Surface quality of the implanted substrate
- Electrical properties, such as carrier mobility, sheet charge concentration, resistivity
- Substitutionality of the implanted dopants

Secondary Ion Mass spectrometry(SIMS)[1] is the preferred technique to evaluate depth profiles of the implanted ions and is used here to evaluate Si and Fe distributions in InP. During SIMS analysis, the sample surface is slowly sputtered away by bombarding with primary ions. The sputtering process produces secondary ions of the constituent elements of the substrate with a range of (translational) kinetic energies. Monitoring the secondary ion count rate of selected elements as a function of time leads to depth profiles. The analysis usually uses  $Cs^+$  or  $O^-$  primary ions. The mass spectra of the secondary ions, obtained by continuously monitoring the ion signal while scanning a range of mass-to-charge ( $m/z$ ) ratios, detects and distinguishes both atomic and molecular ions. The depth resolution of SIMS is between 2 - 30 nm. Mass interferences occur whenever another ion has the same nominal mass as the analyzed ion. Such interferences are called isobaric. During the analysis of iron in silicon for example,  $^{28}Si^{2+}$  interferes because it has the same mass-charge as  $^{56}Fe^+$ . In this case isotopes such as  $^{29}Si$  or  $^{54}Fe$  are detected and scaled using the corresponding sensitivity factors. SIMS is a destructive technique due to the sputtering away of the constituent elements.

## APPENDIX A. IMPLANTATION ANALYSIS

### Instrumental Conditions

<i>Condition Number</i>	1	2
<i>Instrument</i>	Cameca	Cameca
<i>Elements Monitored</i>	Si	Fe
<i>Primary Ion Beam</i>	Cs	Cs
<i>Primary Ion Energy</i>	14.5keV	5.5keV
<i>Secondary Ion Polarity</i>	Negative	Positive
<i>Oxygen Leak</i>	No	No
<i>Charge Neutralization†</i>	No	No
<i>Surface Conductive Coating†</i>	None	None
<i>Liquid Nitrogen Cold Trap*</i>	Yes	No

† Used for accurate analysis of insulating samples or substrates.

\* Used to minimize instrumental background for atmospheric impurities.

Figure A.1: The conditions used for SIMS in this dissertation

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To evaluate the damage or crystallinity, various methods are used such as Rutherford Backscattering spectrometry (RBS), Transmission Electron Microscopy (TEM), X-Ray Diffraction (XRD) and Photoluminescence spectra (PLs). Only RBS is discussed here since this is primarily used for evaluating crystal quality. Rutherford backscattering [2] detects the energies and amount of the backscattered ions from a solid target. Ion channeling is a specific application of RBS in which the absence of backscattering signal is quantified. In an ion backscattering application, alignment of the incident beam with a major crystallographic axis will allow the incident ions to be channeled down the open regions of the crystal lattice. For a channeled orientation, the incident ions can be transported to very great depths ( $\geq 3\text{-}5\ \mu\text{m}$ ) from which backscattering events cannot be detected. If the crystal lattice contains either matrix or impurity atoms which are not in the crystal structure, that is, interstitial in the crystal structure, backscattering events can quantify the concentration of interstitial atoms (in units of atoms/cm<sup>2</sup>). Thus, range and extent of damage from ion implantation, degree of crystal regrowth upon annealing, and the degree of impurity substitutionality/interstitiality can all be quantified in an ion channeling experiment. Here channeled and rotating random RBS spectra are acquired at detector angles of 160° and 100° from the forward trajectory of the incident He ion beam. The different depth resolution and backscattering kinematics afforded by the use of two different angles improve the accuracy of the measurements. The grazing detector spectra are sensitive to the surface layers of the crystal and provide more precise damage



## APPENDIX A. IMPLANTATION ANALYSIS

depths. When the sample is moved in a rotating random orientation, the sample normal is precessed about the incident ion-beam. The backscattering spectrum from this orientation is representative of a spectrum from a polycrystalline sample and is normalizable as the signal for 100% amorphization at all depths. The channeled spectra can then be compared to the rotating random spectra to yield quantifiable channeling parameters.

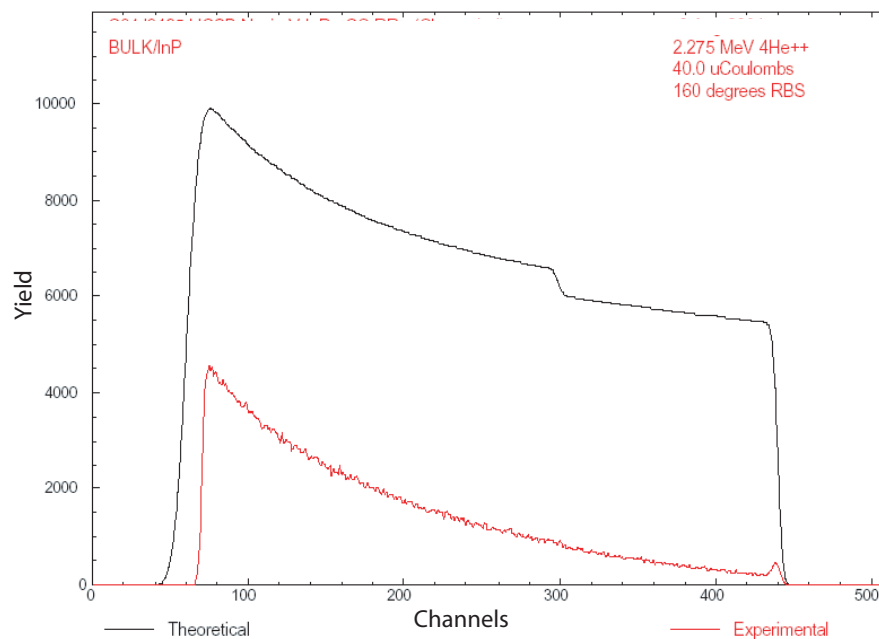


Figure A.2: A sample RBS spectra for SI InP:Fe substrate

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To evaluate surface morphology, Atomic Force Microscopy (AFM) [3] is used. These are variations on a method of imaging surfaces with atomic or near-atomic resolution, collectively called scanning probe microscopy (SPM). A small tip is scanned across the surface of a sample in order to construct a 3-D image of the surface. Fine control of the scan is accomplished using piezoelectrically-induced motions. Any type of surface can be probed by the molecular forces exerted by the surface against the tip. The tip can be constantly in contact with the surface, it can gently tap the surface while oscillating at high frequency, or it can be scanned just minutely above the surface. Image processing software allows easy extraction of useful surface parameters.

## APPENDIX A. IMPLANTATION ANALYSIS

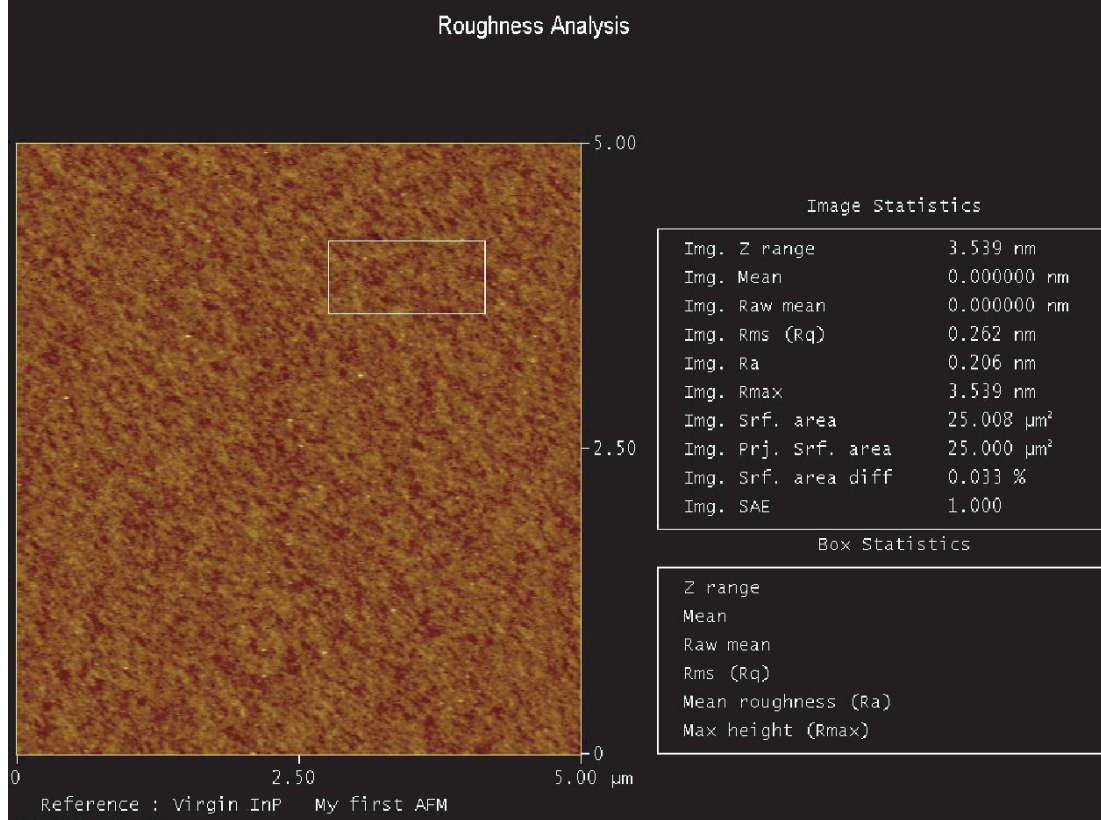


Figure A.3: AFM scan of SI InP substrate

Electrical properties such as mobility and sheet charge density are obtained from Hall measurements. The basic physical principle underlying the Hall effect [4] is the Lorentz force. When an electron current is (due to an applied Electric Field) in a direction perpendicular to an applied magnetic field, it experiences a force acting normal to both directions and moves in response to this force, resulting in an excess surface electrical charge on the side of the sample. This charge results in the transverse Hall voltage that can be measured. The sheet charge density can be extracted as

$$n_s = \frac{I \times B}{q \times |V_H|} \quad (\text{A.0.2})$$

Thus, by measuring the Hall voltage  $V_H$  and from the known values of  $I$ ,  $B$ , and  $q$ , one can determine the sheet density  $n_s$  of charge carriers in semiconductors. The Hall voltage is negative for n-type semiconductors and positive for p-type semi-

## APPENDIX A. IMPLANTATION ANALYSIS

conductors. The sheet resistance  $R_S$  of the semiconductor can be conveniently determined by use of the van der Pauw resistivity measurement technique [5]. Since sheet resistance involves both sheet density and mobility, one can determine the Hall mobility from the equation

$$\mu = \frac{1}{qn_s R_s} \quad (\text{A.0.2})$$

## References

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# B

## Implanted collector InP HBT / Circuit Process Flow

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THIS appendix details the process flow for fabricating implanted collector InP DHBTs and circuits.

- The high temperature annealing steps are critical
  - Clean the quartz chamber in the RTA thoroughly with solvents. Clean the RTA chamber by flowing  $N_2$  gas for at least 1 hour just prior to actual anneal. Do at least 2-3 dummy runs before actual anneal.
  - A very smooth  $Si_xN_y$  is very important for a defect free anneal. Clean the PECVD chamber for at least 1 hour prior to deposition. When doing backside deposition, do not slide the wafer off the platen, and ensure that the platen is coated first with at least 100 nm of  $Si_xN_y$ .
  - Pristine surface quality is to be maintained for all steps prior to growth. The wafer should always be handled at the edges with a clean tweezer. There should be absolutely no scratches on the wafer.
  - Any organics present on the wafer will give rise to pitting at the surface during anneal. The wafer should be very clean of PR and other organics before encapsulation with  $Si_xN_y$ .
- Extra precaution should be exercised when handling a 2 inch InP wafer.
  - Before spinning PR, always check the chuck vacuum during spin with a dummy 2 inch GaAs wafer.
  - When placing clips on the wafer during sputtering or evaporation, make sure that they are not in the same direction from the major and minor flats (or the wafer will crack).

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Sometimes, the PR + a thick  $Si_xN_y$  implant mask will fail to resolve the alignment targets for local alignment in the GCA stepper. In this case, open windows over one set of alignment targets.
- The process tolerances are sensitive to the quality of the photoresist
  - If uncertain of the age and/or quality of the PR, pour a new bottle
  - This is especially important for SPR-518, SPR-510, SPR-955, CEM, and nLOF 2020
- When spinning PR, use a slow acceleration
  - i.e. It should take 2 → 2.5 seconds to reach maximum spin speed
- For the following steps, a focus array is required for accurate features:
  1. Ni alignment targets lithography
  2. Pedestal implant lithography
  3. Emitter contact lithography
  4. Base contact lithography
  5. Base post lithography
  - Otherwise, the focus offset on subsequent steps is = 0
  - Always inspect the focus checkers thoroughly after development
- **Wafer orientation**
  - There are two type of wafers
    1. European / Japan flat option wafer where the minor flat is to the left of the major flat
    2. US flat option wafer where the minor flat is to the right of the major flat
  - To ensure proper semiconductor mesa etch undercut, the long-axis of the emitter should be oriented...
    - \* Perpendicular to the major flat for European / Japan flat option wafers
    - \* Parallel to the major flat for US flat option wafers
  - MBE growth can be performed only on full 2 inch InP wafers. The wafer is cleaved after growth

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

### 1. Blanket Fe implant and anneal

- Ship the 2 inch SI epi-quality InP wafers for Fe implant. For implanted subcollector DHBTs, choose low energy-low dose implant as described in Chapter 4 . For the implanted pedestal-subcollector DHBTs, choose a higher energy (typically  $\sim 150$  keV) and higher fluence ( $1-3 \times 10^{14}$  ions/cm<sup>2</sup>). The implants are always carried out at 7° offset and at 200 °C. This implant temperature is critical as described in Chapter 3 . Implant current should be maintained  $\leq 50 \mu A$ .
- Record  $N_s, K_s$  from ellipsometer
- Deposition of  $Si_xN_y$  in the PECVD
  - Clean insides of PECVD with DI first, then ISO.
  - Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
  - Surface preparation – 20 sec BHF + 2 min DI water rinse
  - Deposit 40 nm of  $Si_xN_y$  at 265 °C on the wafer
  - Deposit 40 nm of  $Si_xN_y$  at 265 °C on the platen
  - Prepare ozone reactor – run empty, 20 min
  - Surface Preparation for  $Si_xN_y$  deposition 1 – Oxidise wafer in uv Ozone for 120 sec
  - Surface Preparation for  $Si_xN_y$  deposition 2 – 30 sec clean in BHF, rinse in DI for 5 min
  - Deposit 40 nm of  $Si_xN_y$  on the top side
  - Solvent clean – 3 min 2-propanol, 3 min DI water rinse
  - Check quality of SiNx
  - If  $Si_xN_y$  looks rough or non uniform: remove in BHF for 10 min and rinse in running DI water for 5-10 min and repeat above steps from the uv Ozone. If  $Si_xN_y$  still looks rough, increase uv Ozone time to 200 sec
  - Deposit 100 nm of  $Si_xN_y$  on PECVD platen
  - Place wafer very, very gently with top side on platen. **Do not let it slide**
  - Deposit 40 nm of  $Si_xN_y$  on the back side
- After 1h clean of RTA, run empty 700 °C for 300s
- Run actual wafer anneal at 700 °C for 300s
- Remove  $Si_xN_y$  cap by agitating in BHF for  $\sim 10$  min

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Rinse in running DI for 5 min
- Inspect under microscope to ensure that is no extensive pitting on the surface
- Record  $N_s$ ,  $K_s$  using P270000 on Ellipsometer. When annealed fully, these should be same as for virgin SI InP substrate

### 2. Alignment targets definition

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- HF clean – 5 min conc. HF:DI (10 ml: 200 ml), 5 min in running DI water
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Sputter TiW
  - Prepare sputter tool – load TiW source, TiW anode, TiW shims and W shield
  - Load wafer and after a quick pumpdown, ensure correct operation by turning on DC power and checking if plasma is light violet-blue
  - Ensure that the system pumps down to  $\leq 3 \times 10^{-7}$  Torr. At  $\leq 1 \times 10^{-6}$  Torr, add  $LN_2$  if necessary
  - RF back sputter for 2 min at 100 V
  - Sputter for 1-2 min at 200W with shutter closed
  - Open shutter and sputter on wafer for 30 min at 200W, for  $\sim 350$  nm of TiW
- Inspect under microscope
- Deposition of  $Si_xN_y$  in the PECVD
  - Clean insides of PECVD with DI first, then ISO.
  - Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
  - Surface preparation – 20 sec BHF + 2 min DI water rinse
  - Deposit 40 nm of  $Si_xN_y$  at 265 °C on the wafer
- Lithography and for Ni metal liftoff
  - Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
  - Photoresist spin – LOL-2000, 1 kRPM, 45 sec

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Pre-exposure PR soft bake – 210 °C, 5 min
- Photoresist spin – SPR-955-0.9CM, 3 kRPM, 30 sec
- Pre-exposure PR bake – 95 °C, 1 min
- Shoot ‘Alignment Mark ’ pattern in stepper, 0.9 sec
- Post-exposure PR bake – 110°C, 60 sec
- Develop in MF-701 developer, 60 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope
- Vent E-beam 4, load private source – Ni
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit 300 Å of Ni at 1 Å/s
- Metal liftoff – 1165 stripper, 80°C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80°C, 20 min
- Gently agitate sample surface with small pipet
- Rinse sample in 2-propanol before transferring to H<sub>2</sub>O or Solvent clean
- If scum present, squirt with pipette and leave in PRX-127 (at  $\geq 90$  °C) for 15 min
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dry etch alignment targets
  - Clean RIE<sub>3</sub> chamber with O<sub>2</sub>/50 mT/500 V for 60 min
  - Dry etch TiW with SF<sub>6</sub>/Ar for 2.5/10 sccm, 20 mT, 100 V – Use laser monitor to determine stop time
  - Over etch for 1 min 30 s
  - Inspect under microscope
  - Check height using profilometer

### 3. Deposition of $Si_xN_y$ as implant mask using the PECVD

The thickness of the  $Si_xN_y$  implant mask is determined by the implant conditions. Use TRIM and the discussion in this thesis to determine the thickness of  $Si_xN_y$ . For 350 keV Si implant, 140 nm of  $Si_xN_y$  is sufficient. Also load a GaAs dummy wafer along with actual wafer during all depositions

- Clean insides of PECVD with DI first, then ISO.
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program



## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Prepare ozone reactor – run empty, 20 min
- Surface Preparation for  $Si_xN_y$  deposition :1 – Oxidise wafer in uv Ozone for 200 sec
- Surface Preparation for  $Si_xN_y$  deposition :2 – 30 sec clean in BHF, rinse in DI for 5 min
- Deposit 40 nm of  $Si_xN_y$  on the wafer
- Check quality of  $Si_xN_y$ . If  $Si_xN_y$  looks rough or non uniform: remove in BHF for 10 min and rinse in DI 5-10 min and repeat above steps from the uv Ozone. If  $Si_xN_y$  still looks rough, increase uv Ozone time to 300 sec
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- If  $Si_xN_y$  is uniform and smooth, deposit 200 nm more of  $Si_xN_y$  on the wafer and the dummy GaAs
- Inspect under microscope
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- If  $Si_xN_y$  is uniform and free of pinholes, deposit 400 nm more of  $Si_xN_y$  on the wafer and the dummy GaAs
- Inspect under microscope
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- If  $Si_xN_y$  is uniform and free of pinholes, deposit 400 nm more of  $Si_xN_y$  on the wafer and the dummy GaAs
- Inspect under microscope
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- If  $Si_xN_y$  is uniform and free of pinholes, deposit 400 nm more of  $Si_xN_y$  on the wafer and the dummy GaAs
- Record  $N_s$ ,  $K_s$  using P270000 on Ellipsometer and calculate thickness from Filmetrics

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

### 4. Subcollector Implant Mask lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-518, 4 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- Shoot ‘Align Window’ pattern in stepper, 0.7 sec – deliberately underexposed to not clear all photoresist over alignment targets
- Mini post-exposure PR bake – 90 °C, 10 sec
- Develop in MF-701 for 40 sec
- Rinse wafer – DI water for 20 sec, N<sub>2</sub> dry
- Inspect wafer using optical microscope and measure PR removed using profilometer
- Second pre-exposure PR bake – 90 °C, 20 sec
- Shoot ‘Subcollector Implant’ pattern in stepper, 1.0 sec
- Post-exposure PR bake – 105 °C, 10 sec
- Develop in MF-701 for 2 min
- Rinse wafer – DI water for 3 min sec, N<sub>2</sub> dry
- Inspect wafer using optical microscope and measure PR height profilometer

### 5. Dry etch of $Si_xN_y$ implant mask

- uv flood expose wafer for 1 - 2 min
- Clean RIE<sub>3</sub> chamber with O<sub>2</sub> at 50 mT, 500 V for 60 min
- Place dummy real wafer and GaAs wafer with the same thickness of  $Si_xN_y$  in the chamber. Ensure that the laser strikes the center of the GaAs dummy wafer.
- Pump down to  $\leq 9 \times 10^{-6}$  T. Dry etch using CF<sub>4</sub> : O<sub>2</sub>, 20 : 2 sccm, 20 mT, 200 V. The etch rate is  $\approx 400$  Å / min
- Monitor reflected laser signal carefully. Note the time at which the etch has stopped, as indicated by a constant intensity of the reflected laser signal. Over etch for 2 min at 200 V, and then for 2 more min at 100 V.

## APPENDIX B. IMPLANTED COLLECTOR IN PHBT / CIRCUIT PROCESS FLOW

- Inspect using optical microscope if  $Si_xN_y$  is removed fully wherever Si is to be implanted
- Measure height of PR + implant mask
- If  $Si_xN_y$  is uniform repeat dry etch with 1 min increments.
- uv flood expose wafer for 1 - 2 min
- Strip PR in conc. AZ-400K for 3 min
- Gently agitate sample surface with small pipet
- Rinse wafer – DI water for 2 min,  $N_2$  dry
- If scum present, squirt with pipette and leave in PRX-127 (at  $\geq 90$  °C) for 15 min
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean
- Inspect wafer using optical microscope and measure thickness of  $Si_xN_y$  implant mask with the profilometer
- Carefully record misalignment of the subcollector implant pattern with the zero level targets over the entire wafer
- SEM subcollector implant patterns to measure actual dimensions of the subcollector implant windows
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min

### 6. Ship for Si subcollector implant

For implanted subcollector-pedestal implant, a multiple energy-fluence implant is performed to obtain  $\sim 1 \times 10^{19} \text{ cm}^{-3}$  over  $\sim 400 - 500 \text{ nm}$  for a low sheet resistance of the subcollector and a high surface doping. For implanted pedestal - subcollector process, Si implant is performed at 350 keV with a fluence of  $\leq 6 \times 10^{14} \text{ ions/cm}^2$ . Chapter 3 details the implant process.

### 7. Post implant processing – $Si_xN_y$ (and Ni) removal

- Inspect wafers in the optical microscope and record misalignment again
- If scum present, squirt with pipette and leave in PRX-127 (at  $\geq 90$  °C) for 15 min
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean
- Prepare HF beakers – rinse thoroughly in DI and then briefly in BHF

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Agitate in 250 ml of HF continuously till  $Si_xN_y$  is seen to disappear and the wafer becomes colorless. Over etch for 10 min and ensure that no  $Si_xN_y$  remains
- Try to measure  $N_L, T_L$  of  $Si_xN_y$  on Ellipsometer – if  $Si_xN_y$  is fully gone, the reading will be in error
- Inspect wafers in the optical microscope and ensure that  $Si_xN_y$  is completely removed Also, ensure that Ni and  $Si_xN_y$  on top of TiW alignment targets are completely removed
- if Ni is not fully removed, repeat BHF wet etch with constant agitation in 5 min increments
- Measure height of alignment targets profilometer. Check profile across the implanted regions – there should be less than  $\sim 50 \text{ \AA}$  variation

### 8. High temperature anneal

- Record  $N_s, K_s$  of the InP wafer using P270000 on Ellipsometer
- Deposition of  $Si_xN_y$  in the PECVD
  - Clean insides of PECVD with DI first, then ISO.
  - Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
  - Surface preparation – 20 sec BHF + 2 min DI water rinse
  - Deposit 40 nm of  $Si_xN_y$  at 265 °C on the wafer
  - Deposit 40 nm of  $Si_xN_y$  at 265 °C on the platen
  - Prepare ozone reactor – run empty, 20 min
  - Surface Preparation for  $Si_xN_y$  deposition 1 – Oxidise wafer in uv Ozone for 100 sec
  - Surface Preparation for  $Si_xN_y$  deposition 2 – 30 sec clean in BHF, rinse in DI for 5 min
  - Deposit 40 nm of  $Si_xN_y$  on the top side
  - Solvent clean – 3 min 2-propanol, 3 min DI water rinse
  - Check quality of SiNx
  - If  $Si_xN_y$  looks rough or non uniform: remove in BHF for 10 min and rinse in running DI water for 5-10 min and repeat above steps from the uv Ozone. If  $Si_xN_y$  still looks rough, increase uv Ozone time to 150 sec
  - Deposit 100 nm of  $Si_xN_y$  on PECVD platen

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Place wafer very, very gently with top side on platen. **Do not let it slide**
- Deposit 40 nm of  $Si_xN_y$  on the back side
- After 1h clean of RTA, run empty 800 °C for 30 sec, twice
- For implanted subcollector process, run actual wafer anneal at 800 °C for 30 sec  
For implanted subcollector-pedestal process, run actual wafer anneal at 600 °C for 10 sec
- Remove  $Si_xN_y$  cap by agitating in BHF for ~ 10 min
- Rinse in running DI for 5 min
- Inspect under microscope to ensure that is no extensive pitting on the surface
- Record  $N_s, K_s$  of the wafer using P270000 on Ellipsometer. When annealed fully, these should be same as for virgin SI InP substrate

For a implanted subcollector process, proceed straight to surface preparation in step 15.

For implanted subcollector-pedestal process, proceed to next step which is definition of  $Si_xN_y$  mask for pedestal implant

### 9. Deposition of $Si_xN_y$ as pedestal implant mask using the PECVD

The thickness of the  $Si_xN_y$  implant mask is determined by the implant conditions. Use TRIM and the discussion in this thesis to determine the thickness of  $Si_xN_y$ . For 100 keV Si implant, 80 nm of  $Si_xN_y$  is sufficient. Also load a GaAs dummy wafer along with actual wafer during all depositions

- Clean insides of PECVD with DI first, then ISO.
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
- Prepare ozone reactor – run empty, 20 min
- Surface Preparation for  $Si_xN_y$  deposition :1 – Oxidise wafer in uv Ozone for 200 sec
- Surface Preparation for  $Si_xN_y$  deposition :2 – 30 sec clean in BHF, rinse in DI for 5 min
- Deposit 40 nm of  $Si_xN_y$  on the wafer

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Check quality of SiNx. If  $Si_xN_y$  looks rough or non uniform: remove in BHF for 10 min and rinse in DI 5-10 min and repeat above steps from the uv Ozone. If  $Si_xN_y$  still looks rough, increase uv Ozone time to 300 sec
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- If  $Si_xN_y$  is uniform and smooth, deposit 200 nm more of  $Si_xN_y$  on the wafer and the dummy GaAs
- Inspect under microscope
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- If  $Si_xN_y$  is uniform and free of pinholes, deposit 300 nm more of  $Si_xN_y$  on the wafer and the dummy GaAs
- Inspect under microscope
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- If  $Si_xN_y$  is uniform and free of pinholes, deposit 400 nm more of  $Si_xN_y$  on the wafer and the dummy GaAs
- Inspect under microscope
- Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- Record  $N_s, K_s$  using P270000 on Ellipsometer and calculate thickness from Filmetrics

### 10. Pedestal Implant lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-510, 3 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- Shoot ‘Pedestal Implant’ pattern in stepper, 1.4 sec

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Post-exposure PR bake – 105 °C, 10 sec
- Develop in MF-701 for 2 min
- Rinse wafer – DI water for 3 min sec, N<sub>2</sub> dry
- Inspect wafer using optical microscope. If misalignment to ZLTs is more than 0.1 μm, repeat lithography

### 11. Dry etch of $Si_xN_y$ implant mask

- uv flood expose wafer for 1 - 2 min
- Clean RIE<sub>3</sub> chamber with O<sub>2</sub> at 50 mT, 500 V for 60 min
- Place dummy real wafer and GaAs wafer with the same thickness of  $Si_xN_y$  in the chamber. Ensure that the laser strikes the center of the GaAs dummy wafer.
- Pump down to  $\leq 9 \times 10^{-6}$  T. Dry etch using CF<sub>4</sub> : O<sub>2</sub>, 20 : 2 sccm, **10 mT**, 200 V. The etch rate is  $\approx 400 \text{ \AA} / \text{min}$
- Monitor reflected laser signal carefully. Note the time at which the etch has stopped, as indicated by a constant intensity of the reflected laser signal. Over etch for 2 min at 200 V, and then for 2 more min at 100 V.
- Inspect using optical microscope if  $Si_xN_y$  is removed fully wherever Si is to be implanted
- Measure height of PR + implant mask
- If  $Si_xN_y$  is uniform repeat dry etch with 1 min increments.
- uv flood expose wafer for 1 - 2 min
- Strip PR in conc. AZ-400K for 3 min
- Gently agitate sample surface with small pipet
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- If scum present, squirt with pipette and leave in PRX-127 (at  $\geq 90$  °C) for 15 min
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean
- Inspect wafer using optical microscope and measure thickness of  $Si_xN_y$  implant mask with the profilometer
- Carefully record misalignment of the pedestal implant pattern with the zero level targets over the entire wafer

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- SEM pedestal implant patterns to measure actual dimensions of the window
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min

### 12. Ship for Si pedestal implant

For implanted subcollector-pedestal implant, a multiple energy-fluence implant is performed to obtain  $\sim 1 \times 10^{19} \text{ cm}^{-3}$  over the existing Fe doping. Chapter 5 details the implant conditions.

### 13. Post implant processing – $Si_xN_y$ (and Ni) removal

- Inspect wafers in the optical microscope and record misalignment again
- If scum present, squirt with pipette and leave in PRX-127 (at  $\geq 90^\circ\text{C}$ ) for 15 min
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean
- Prepare HF beakers – rinse thoroughly in DI and then briefly in BHF
- Agitate in 250 ml of HF continuously till  $Si_xN_y$  is seen to disappear and the wafer becomes colorless. Over etch for 10 min and ensure that no  $Si_xN_y$  remains
- Try to measure  $N_L, T_L$  of  $Si_xN_y$  on Ellipsometer – if  $Si_xN_y$  is fully gone, the reading will be in error
- Inspect wafers in the optical microscope and ensure that  $Si_xN_y$  is completely removed.
- Measure height of alignment targets profilometer. Check profile across the implanted regions – there should be less than  $\sim 50 \text{ \AA}$  variation

### 14. High temperature anneal

- Record  $N_s, K_s$  of the InP wafer using P270000 on Ellipsometer
- Deposition of  $Si_xN_y$  in the PECVD
  - Clean insides of PECVD with DI first, then ISO.
  - Perform 30 minutes clean of the PECVD and condition with  $Si_xN_y$ – use standard program
  - Surface preparation – 20 sec BHF + 2 min DI water rinse
  - Deposit 40 nm of  $Si_xN_y$  at 265 °C on the wafer



## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Deposit 40 nm of  $Si_xN_y$  at 265 °C on the platen
- Prepare ozone reactor – run empty, 20 min
- Surface Preparation for  $Si_xN_y$  deposition 1 – Oxidise wafer in uv Ozone for 100 sec
- Surface Preparation for  $Si_xN_y$  deposition 2 – 30 sec clean in BHF, rinse in DI for 5 min
- Deposit 40 nm of  $Si_xN_y$  on the top side
- Solvent clean – 3 min 2-propanol, 3 min DI water rinse
- Check quality of SiNx
- If  $Si_xN_y$  looks rough or non uniform: remove in BHF for 10 min and rinse in running DI water for 5-10 min and repeat above steps from the uv Ozone. If  $Si_xN_y$  still looks rough, increase uv Ozone time to 150 sec
- Deposit 100 nm of  $Si_xN_y$  on PECVD platen
- Place wafer very, very gently with top side on platen. **Do not let it slide**
- Deposit 40 nm of  $Si_xN_y$  on the back side
- After 1h clean of RTA, run empty 800 °C for 30 sec, twice
- Run actual wafer anneal at 800 °C for 30 sec
- Remove  $Si_xN_y$  cap by agitating in BHF for ~ 10 min
- Rinse in running DI for 5 min
- Inspect under microscope to ensure that is no extensive pitting on the surface
- Record  $N_s$ ,  $K_s$  of the wafer using P270000 on Ellipsometer. When annealed fully, these should be same as for virgin SI InP substrate

If the alignment targets are stable, the wafers are ready for growth.

### 15. Surface preparation

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Prepare ozone reactor – run empty, 20 min
- Surface preparation 1 – oxidize wafer surface in ozone reactor, 10 min
- Surface preparation 2 – conc. HF:DI water 5 ml:200 ml for 5 min
- Surface preparation 3 – Rinse in running DI water for 5 min
- Load sample **immediately** in the MBE chamber

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**MBE growth of drift collector, base, emitter layers. RHEED is continuously monitored to ensure that the growth is excellent**

**16. Wafer preparation for emitter lithography – wet etch of semiconductor over TiW alignment targets**

- Inspect wafer in the SEM
- Inspect wafer under the optical microscope. For the GCS stepper, alignment marks after regrowth are not resolved by the DFAS local alignment system. The layers on top of the alignment targets have to be removed
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-510, 4 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- Shoot ‘Align window’ pattern in stepper, 1.4 sec
- Post-exposure PR bake – 105 °C, 10 sec
- Develop in MF-701 for 2 min
- Rinse wafer – DI water for 3 min sec, N<sub>2</sub> dry
- Inspect wafer using optical microscope
- Prepare two beakers with...
  - (a) H<sub>3</sub>PO<sub>4</sub>:DI water<sub>2</sub>:DI water, 1:1:25 – use stirrer at 200 RPM
  - (b) H<sub>3</sub>PO<sub>4</sub>:HCl, 4:1 – use stirrer at 200 RPM
- Etch InGaAs emitter cap in H<sub>3</sub>PO<sub>4</sub>:H<sub>2</sub>O<sub>2</sub>:DI water ≈ 25 sec
- Inspect under microscope and check in profilometer to ensure all InGaAs is etched
- Etch InP emitter in H<sub>3</sub>PO<sub>4</sub>:HCl ≈ 40 sec
- Inspect under microscope and check in profilometer to ensure all InP is etched
- Etch InGaAs base in H<sub>3</sub>PO<sub>4</sub>:DI water<sub>2</sub>:DI water ≈ 40 sec
- Inspect under microscope and check in profilometer to ensure all InGaAs is etched
- PR removal

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- Place sample in 1165 PR stripper, 80°C, 1 hour
- Gently agitate sample surface with small pipet
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean
- If scum is present, squirt gently in AZ-300T PR stripper, 80°C, 15 min
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse

### 17. Wafer cleaving

- For the local-alignment DFAS (Dark-Field-Alignment-System) system to work on the i-line GCA stepper, the minimum size of the sample to be processed should be greater than 1 x 1 inch<sup>2</sup>.
- Cleave into 4 quarters carefully along the boundary of a die.

### 18. Emitter contact lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-955, 2.5 kRPM, 30 sec
- Pre-exposure PR bake – 92 °C, 60 sec
- CEM coat and spin
  - Coat wafer w/ CEM and let sit for 60 sec
  - Then spin 4 kRPM, 30 sec
- Shoot ‘emitter’ pattern in stepper, 2.055 sec
- Post-exposure PR bake – 110 °C, 60 sec
- Development
  - (a) Rinse CEM from wafer surface using DI water, 30 sec
  - (b) Immediately transfer wafer to MF-701 developer
    - Develop for 2 min 20 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope. Misalignment to pedestal should be ≤ 0.05 μm

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### 19. Emitter contact deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 10 min
- Vent E-beam 4, load private sources – Ti, Pd, and Au
- Surface preparation – HCl:DI water 1:10 dip 10 sec, DI water rinse 10 sec, N<sub>2</sub> dry
- Load sample in E-beam 4 – orient long-axis of emitter in the same direction as the sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit emitter contact
  - Ti 200 Å (1 Å/sec)
  - Pd 400 Å (1 Å/sec)
  - Au 7500 Å
    - \* 1 Å/sec for 1-300 Å
    - \* 2 Å/sec for 301-500 Å
    - \* 3 Å/sec for 501-1000 Å
    - \* 4-5 Å/sec 1001-7500 Å
- Metal liftoff – 1165 stripper, 80 °C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Anneal emitter contact, 300 °C, N<sub>2</sub> purge, 60 sec

### 20. Emitter mesa etch

- Prepare three beakers with...
  - (a) NH<sub>4</sub>OH:DI water, 1:10
  - (b) H<sub>3</sub>PO<sub>4</sub>:H<sub>2</sub>O<sub>2</sub>:DI water, 1:1:25 – use stirrer at 200 RPM
  - (c) H<sub>3</sub>PO<sub>4</sub>:HCl, 4:1 – use stirrer at 200 RPM
- Dip sample – NH<sub>4</sub>OH:DI water solution 10 sec, DI water rinse 10 sec, N<sub>2</sub> dry

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- Etch InGaAs emitter cap in  $\text{H}_3\text{PO}_4:\text{H}_2\text{O}_2:\text{DI water} \approx 21$  sec
  - Overetch should only be 3 sec from the time the color change is complete
  - Inspect under microscope to ensure all InGaAs is etched
- Etch InP emitter in  $\text{H}_3\text{PO}_4:\text{HCl} \approx 35$  sec
  - Overetch should only be 5 sec from the time the color change is complete
  - Inspect under microscope to ensure all InP is etched

### 21. Base contact lithography

- No solvent cleaned needed after emitter mesa etch
- Dehydration bake –  $120^\circ\text{C}$ , 10 min
- Photoresist spin – nLOF 2020, 4 kRPM, 30 sec
- Pre-exposure PR bake –  $111^\circ\text{C}$ , 60 sec
- Shoot ‘base contact’ pattern in stepper – 0.46 sec
- Post-exposure PR bake –  $114^\circ\text{C}$ , 60 sec
- Development – MF-701 developer, 2 min
- Rinse wafer – DI water for 2 min,  $\text{N}_2$  dry
- Inspect wafer using optical microscope

### 22. Base contact deposition

- Prepare ozone reactor – let run empty for 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 10 min
- Vent E-beam 4, load private sources – Ti, Pd, and Au
- Surface preparation –  $\text{NH}_4\text{OH}:\text{DI water } 1:10$  dip 10 sec,  $\text{N}_2$  dry, **NO WATER RINSE**
- Load sample in E-beam 4 – orient long-axis of emitter in the same direction as the sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit base contact
  - Pd 25 Å, Ti 170 Å, Pd 170 Å, Au 650 Å
  - \* 1 Å/sec for all metal depositions

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- \* Higher deposition rates onto nLOF 2020 will leave more PR scum
- Metal liftoff – 1165 stripper, 80°C, 1 hour
- Remove metal and place sample in fresh AZ 300T stripper, 80°C, 10 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean
  - The sample can proceed to ‘base-post’ lithography if there is no scum on post-end of the base contact
- \* Otherwise, repeat the AZ 300T step until that is the case

### 23. Base post lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-955, 2.5 kRPM, 30 sec
- Pre-exposure PR bake – 92°C, 60 sec
- CEM coat and spin
  - Coat wafer w/ CEM and let sit for 60 sec
  - Then spin 4 kRPM, 30 sec
- Shoot ‘base post’ pattern in stepper, 2.055 sec
- Post-exposure PR bake – 110°C, 60 sec
- Development
  - (a) Rinse CEM from wafer surface using DI water, 30 sec
  - (b) Immediately transfer wafer to MF-701 developer, develop 2 min 20 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 24. Base post deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 10 min
- Vent E-beam 4, load private sources – Ti, Pd, and Au

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- Surface preparation – HCl:DI water 1:10 dip 10 sec, DI water rinse 10 sec, N<sub>2</sub> dry
- Load sample in E-beam 4 – orient long-axis of emitter in the same direction as the sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit base post
  - Pd 25 Å (1 Å/sec)
  - Ti 170 Å (1 Å/sec)
  - Pd 170 Å (1 Å/sec)
  - Au 9300 Å
    - \* 1 Å/sec for 1-300 Å
    - \* 2 Å/sec for 301-500 Å
    - \* 3 Å/sec for 501-1000 Å
    - \* 4-5 Å/sec 1001-9300 Å
- Metal liftoff – 1165 stripper, 80°C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80°C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 25. Base mesa lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Gently pipette SPR-510 onto the wafer and let it sit for 20-30 sec
- Photoresist spin – SPR-510, 4 kRPM, 30 sec
- Pre-exposure PR bake – 90°C, 60 sec
- Shoot ‘base mesa’ pattern in stepper, 1.0 sec
- Post-exposure PR bake – 110°C, 60 sec
- Development – MF-701 developer, 90 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

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### 26. Base mesa etch

- Prepare three beakers with...
  - (a)  $\text{NH}_4\text{OH}$ :DI water, 1:10
  - (b)  $\text{H}_3\text{PO}_4$ : $\text{H}_2\text{O}_2$ :DI water, 1:1:25 – use stirrer at 200 RPM
  - (c)  $\text{H}_3\text{PO}_4$ :HCl, 4:1 – use stirrer at 200 RPM
- Dip sample –  $\text{NH}_4\text{OH}$ :DI water solution 10 sec, DI water rinse 10 sec,  $\text{N}_2$  dry
- Etch InGaAs base, InGaAs collector setback, and ternary grade in  $\text{H}_3\text{PO}_4$ : $\text{H}_2\text{O}_2$ :DI water  $\approx$  35 sec
  - Overetch should only be 5 sec from the time the color change is complete
  - Inspect under microscope to ensure all InGaAs is etched
- Etch InP collector in  $\text{H}_3\text{PO}_4$ :HCl  $\approx$  35 sec
  - Overetch should only be 10 sec from the time the color change is complete
  - This extended overetch is for increased collector semiconductor undercut
  - Inspect under microscope to ensure all InP is etched
- Strip PR mask – 1165 stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 27. NiCr Resistor lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120°C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-510, 4 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- CEM coat and spin
  - Coat wafer w/ CEM and let sit for 60 sec
  - Then spin 4 kRPM, 30 sec
- Shoot ‘Resistor’ pattern in stepper, 2.2 sec



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- Post-exposure PR bake – 110 °C, 60 sec
- Development
  - (a) Rinse CEM from wafer surface using DI water, 30 sec
  - (b) Immediately thereafter transfer wafer to MF-701 developer, develop 2 min 30 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 28. NiCr Resistor deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 90 sec
- Vent E-beam 1, load private sources – Ti, SiO<sub>2</sub>, NiCr
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Load sample in E-beam 1
- Allow system to pump-down for 60 min to  $< 2 \cdot 10^{-6}$  torr
- Deposit NiCr resistors
  - Ti 50 Å (1 Å/sec)
  - SiO<sub>2</sub> 200 Å (1 Å/sec)
  - NiCr xxx Å (1 Å/sec)
    - \* For NiCr, Cr out-diffuses more quickly than Ni
    - \* Thus, the source resistivity goes down after each deposition
    - \* Refer to the NiCr worksheet for accurate determination of NiCr deposition thickness for 50 Ω/□
- Metal liftoff – 1165 stripper, 80 °C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 29. Collector contact lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120 °C, 10 min

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- Photoresist spin – nLOF 2020, 3.5 kRPM, 30 sec
- Pre-exposure PR bake – 111°C, 60 sec
- Shoot ‘collector contact’ pattern in stepper – 0.46 sec
- Post-exposure PR bake – 114°C, 60 sec
- Development – MF-701 developer, 2 min
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 30. Collector contact deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 90 sec
  - Do not run sample any longer in the ozone reactor
  - Longer ashing will remove PR from above the emitter contact *and* oxidize too much of the InGaAs sub-collector contact layer
- Vent E-beam 4, load private sources – Ti, Pd, and Au
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Load sample in E-beam 4 – orient long-axis of emitter in the same direction as the sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit collector contact
  - Ti 200 Å (1 Å/sec)
  - Pd 400 Å (1 Å/sec)
  - Au 4500 Å
    - \* 1 Å/sec for 1-300 Å
    - \* 2 Å/sec for 301-500 Å
    - \* 3 Å/sec for 501-1000 Å
    - \* 4 Å/sec 1001-4500 Å
- Metal liftoff – 1165 stripper, 80 °C, 2 hour
- Remove metal and gently agitate in fresh AZ 300T stripper, 80 °C, 10 min
- Rinse in 2-propanol before transferring to DI water or Solvent clean

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### 31. Collector post lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake 120 °C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-518, 3.0 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- CEM coat and spin – coat wafer w/ CEM and let sit for 60 sec, then spin 4 kRPM, 30 sec
- Shoot ‘collector post’ pattern in stepper, 2.6 sec
- Post-exposure PR bake – 110 °C, 60 sec
- Development
  - (a) Rinse CEM from wafer surface using DI water, 30 sec
  - (b) Immediately transfer wafer to MF-701 developer, develop 2 min 30 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 32. Collector post deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 5 min
- Vent E-beam 4, load private sources – Ti and Au
- Surface preparation – HCl:DI water 1:10 dip 10 sec, DI water rinse 10 sec, N<sub>2</sub> dry
- Load sample in E-beam 4 – orient long-axis of emitter perpendicular to the direction of sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit collector post
  - Ti 100 Å (1 Å/sec)
  - Au xxxx Å (see list below for deposition rate)
    - \* The collector post needs to be level with the top of the emitter contact
    - \* Dektak to determine Au deposition thickness

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- \* 1 Å/sec for 1-300 Å
- \* 2 Å/sec for 301-500 Å
- \* 3 Å/sec for 501-1000 Å
- \* 4-5 Å/sec 1001-xxxx Å
- Ti 100 Å (1 Å/sec)
- Metal liftoff – 1165 stripper, 80 °C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 33. Resistor post lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake 120 °C, 10 min
- Cool wafer, 5 min
- Photoresist spin – SPR-518, 3.0 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- CEM coat and spin – coat wafer w/ CEM and let sit for 60 sec, then spin 4 kRPM, 30 sec
- Shoot ‘resistor post’ pattern in stepper, 2.6 sec
- Post-exposure PR bake – 110 °C, 60 sec
- Development
  - (a) Rinse CEM from wafer surface using DI water, 30 sec
  - (b) Immediately transfer wafer to MF-701 developer, develop 2 min 30 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 34. Resistor post deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 5 min
- Vent E-beam 4, load private sources – Ti and Au

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- Surface preparation – HCl:DI water 1:10 dip 10 sec, DI water rinse 10 sec, N<sub>2</sub> dry
- Load sample in E-beam 4 – orient long-axis of emitter perpendicular to the direction of sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit resistor post contact
  - Ti 100 Å (1 Å/sec)
  - Au xxxx Å (see list below for deposition rate)
    - \* The resistor post needs to level with the top of the emitter contact
    - \* Dektak to determine Au deposition thickness
    - \* 1 Å/sec for 1-300 Å
    - \* 2 Å/sec for 301-500 Å
    - \* 3 Å/sec for 501-1000 Å
    - \* 4-5 Å/sec 1001-xxxx Å
  - Ti 100 Å (1 Å/sec)
- Metal liftoff – 1165 stripper, 80 °C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 35. BCB passivation

- The ‘Blue Oven’ must be at room temperature 25°C or less before beginning
  - Otherwise, BCB will bubble during the cure and the sample will be ruined
- NOTE: there must not be any stops in the steps from sample surface preparation to loading the spun sample w/ BCB into the Blue oven
  - Any delays will allow increased surface oxide to regenerate on the semiconductor, increasing leakage currents
  - Oxygen contaminates BCB and prolonged exposure will compromise the cure and ruin the sample
- Prepare the ‘Blue Oven’ – run N<sub>2</sub> through chamber at 100%

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- Prepare ozone reactor – run empty, 20 min
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120 °C, 10 min
- Cool wafer, 5 min
- Surface preparation – oxidize wafer surface in ozone reactor, 10 min
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Coat wafer with BCB 3022-35 – let sit on surface for 30 sec
- Spin BCB 1.5 kRPM, 30 sec – BCB thickness  $\approx 1.88 \mu\text{m}$
- Place sample into Al cup holder
  - The holder should be deformed such that the sample is completely level, yet has minimal contact with the bottom holder surface
  - Otherwise, cured BCB that has crept onto the bottom surface may prevent the sample from being removed from the holder
- Place sample (in holder) into the ‘Blue Oven’
- Reduce N<sub>2</sub> flow to 60% after 3 min
- Load and run Program 5 (confirm in case it has been altered)
- Program sequence:
  - (a) 5 min ramp to 50 °C, 5 min soak
  - (b) 15 min ramp to 100 °C, 15 min soak
  - (c) 15 min ramp to 150 °C, 15 min soak
  - (d) 60 min ramp to 250 °C, 60 min soak
  - (e) Natural cool down
  - (f) Oven off
- Remove sample and inspect under the microscope
- Turn off the ‘Blue Oven’

### 36. Wafer planarization and interconnect surface preparation

- Clean carrier wafer in Panasonic ICP – CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 7 min
  - (a) Load sample on carrier wafer, BCB ICP etch CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 2 min
  - (b) Inspect sample in FEI SEM to see if the device contacts and interconnect posts are exposed

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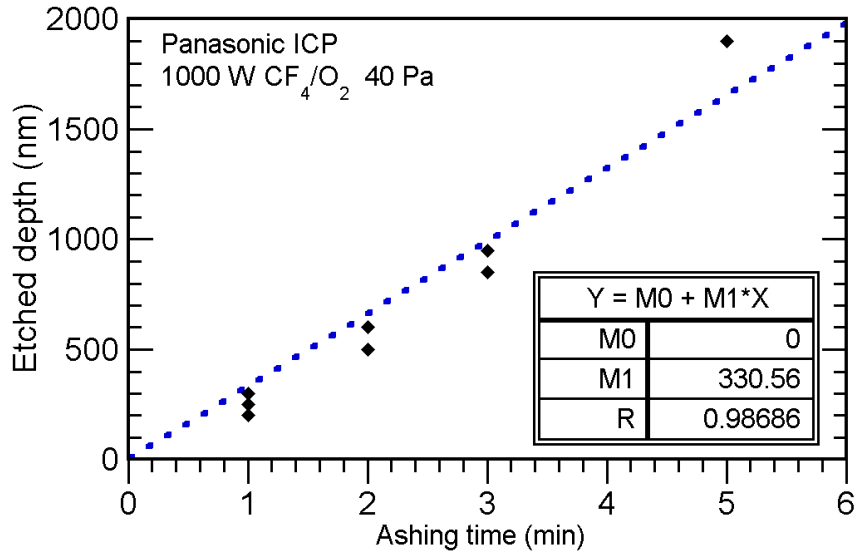


Figure B.1: BCB etch rate for the given CF<sub>4</sub>/O<sub>2</sub> recipe

- Repeat the above etch (1 min increments) and inspection until all contact and posts are exposed
  - Be sure to run the wafer clean program in between each etch cycle
- Repeat the BCB passivation step for further sample planarization
- Clean carrier wafer in Panasonic ICP – CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 7 min
  - (a) Load sample on carrier wafer, BCB ICP etch CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 3 min 30 sec
  - (b) Inspect sample in FEI SEM to see if the device contacts and interconnect post are exposed
- Repeat the above etch (1 min increments) and inspection until all contact and posts are exposed
  - Be sure to run the wafer clean program in between each etch cycle
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake 120 °C, 10 min
- Cool wafer, 5 min
- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 10 min

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- Surface preparation –  $\text{NH}_4\text{OH}$ :DI water 1:10 dip 10 sec,  $\text{N}_2$  dry, **NO WATER RINSE**
- Deposit 100 nm  $\text{SiN}_x$  on the BCB surface by PECVD
- Photoresist spin – SPR-510, 4 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- Shoot ‘contact via’ pattern in stepper, 2.0 sec
- Post-exposure PR bake – 110 °C, 60 sec
- Development – MF-701 developer, 90 sec
- Rinse wafer – DI water for 2 min,  $\text{N}_2$  dry
- Inspect wafer using optical microscope
- Clean carrier wafer in Panasonic ICP –  $\text{CF}_4/\text{O}_2$  50:200 sccm, 7 min
- Load sample on carrier wafer for ICP etching
  - (a)  $\text{SiN}_x$  ICP etch  $\text{CF}_4$  125 sccm, 90 sec
  - (b) Short BCB ashing  $\text{CF}_4/\text{O}_2$  50:200 sccm, 10 sec
- Inspect sample in FEI SEM to see if the device contacts and interconnect posts are exposed
- Strip photoresist mask
  - (a) Flood expose sample, 15 sec
  - (b) Develop exposed PR – MF-701, 2 min
  - (c) Strip remaining PR in acetone – 3 min
  - (d) Inspect – if PR remaining, use 1165 stripper, 80 °C, 3 min
  - (e) Gently agitate sample surface with small pipet
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 37. Metal-1 interconnect lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120 °C, 10 min
- Photoresist spin – nLOF 2020, 4 kRPM, 30 sec
- Pre-exposure PR bake – 111°C, 60 sec
- Shoot ‘Metal 1’ pattern in stepper – 0.46 sec
- Post-exposure PR bake – 114°C, 60 sec



## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Development – MF-701 developer, 2 min
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 38. Metal-1 interconnect deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – clean wafer surface in ozone reactor, 5 min
- Vent E-beam 4, load private sources – Ti and Au
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Load sample in E-beam 4 – orient long-axis of emitter perpendicular to the direction of sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit Metal-1 interconnect
  - Ti 100 Å (1 Å/sec)
  - Au 10 kÅ
    - \* 2 Å/sec for 1-300 Å
    - \* 3 Å/sec for 301-500 Å
    - \* 4 Å/sec for 501-1000 Å
    - \* 5-6 Å/sec 1001-10000 Å
  - Ti 100 Å (1 Å/sec)
- Metal liftoff – AZ 300T stripper, 80 °C, 20 min
- Remove metal and place sample in fresh AZ 300T stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 39. Measure devices and TLMs

- If device performance and TLMs are well behaved, continue

### 40. SiN MIM capacitor formation

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Dehydration bake 120 °C, 10 min
- Cool wafer, 5 min
- Prepare ozone reactor – run empty, 20 min
- Surface preparation – oxidize wafer surface in ozone reactor, 10 min
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Deposit 400 nm SiN<sub>x</sub> onto the sample by PECVD
- Photoresist spin – SPR-510, 4 kRPM, 30 sec
- Pre-exposure PR bake – 90 °C, 60 sec
- Shoot ‘SiN cap’ pattern in stepper, 2.0 sec
- Post-exposure PR bake – 110 °C, 60 sec
- Development – MF-701 developer, 90 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope
- Clean carrier wafer in Panasonic ICP – CF<sub>4</sub>/O<sub>2</sub> 200:40 sccm, 7 min
- Load sample on carrier wafer for ICP etching
  - (a) SiN<sub>x</sub> ICP etch CF<sub>4</sub> 125 sccm, 3 min
  - (b) Short ashing CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 15 sec
- Strip photoresist mask
  - (a) Flood expose sample, 15 sec
  - (b) Develop exposed PR – MF-701, 2 min
  - (c) Strip remaining PR in acetone – 3 min
  - (d) Inspect – if PR remaining, use 1165 stripper, 80 °C, 3 min
  - (e) Gently agitate sample surface with small pipet
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 41. Metal-2 interconnect lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120 °C, 10 min
- Photoresist spin – nLOF 2020, 3 kRPM, 30 sec
- Pre-exposure PR bake – 111 °C, 60 sec

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Shoot 'Metal 2' pattern in stepper – 0.46 sec
- Post-exposure PR bake – 114°C, 60 sec
- Development – MF-701 developer, 2 min
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 42. Metal-2 interconnect deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – clean wafer surface in ozone reactor, 5 min
- Vent E-beam 4, load private sources – Ti and Au
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Load sample in E-beam 4 – orient long-axis of emitter perpendicular to the direction of sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to < 10<sup>-6</sup> torr
- Deposit Metal-2 interconnect
  - Ti 100 Å (1 Å/sec)
  - Au 10 kÅ
    - \* 2 Å/sec for 1-300 Å
    - \* 3 Å/sec for 301-500 Å
    - \* 4 Å/sec for 501-1000 Å
    - \* 5-6 Å/sec 1001-10000 Å
  - Ti 100 Å (1 Å/sec)
- Metal liftoff – 1165 stripper, 80 °C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 43. Metal-2 to Metal-3 interconnect post lithography – 1

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120 °C, 10 min
- Photoresist spin – SPR 518, 3 kRPM, 30 sec

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Pre-exposure PR bake – 90 °C, 60 sec
- CEM coat and spin
  - Coat wafer w/ CEM and let sit for 60 sec
  - Then spin 4 kRPM, 30 sec
- Shoot ‘M2 - M3 post’ pattern in stepper – 2.8 sec
- Post-exposure PR bake – 110 °C, 60 sec
- Development – MF-701 developer, 2 min 30 sec
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 44. Metal-2 to Metal-3 interconnect post deposition – 1

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – clean wafer surface in ozone reactor, 5 min
- Vent E-beam 4, load private sources – Ti and Au
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Load sample in E-beam 4 – orient long-axis of emitter perpendicular to the direction of sample rotation inside E-beam 4
- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit M2-M3 post contact
  - Ti 100 Å (1 Å/sec)
  - Au 13 kÅ
    - \* 2 Å/sec for 1-300 Å
    - \* 3 Å/sec for 301-500 Å
    - \* 4 Å/sec for 501-1000 Å
    - \* 5-6 Å/sec 1001-13000 Å
  - Ti 100 Å (1 Å/sec)
- Metal liftoff – 1165 stripper, 80 °C, 30 min
- Remove metal and place sample in fresh 1165 stripper, 80 °C, 10 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

### 45. Thin-film BCB microstrip wiring environment – layer 1

- The ‘Blue Oven’ must be at room temperature 25°C or less before beginning
  - Otherwise, BCB will bubble during the cure and the sample will be ruined
- NOTE: there must not be any stops in the steps from sample surface preparation to loading the spun sample w/ BCB into the Blue oven
  - Any delays will allow increased surface oxide to regenerate on the surface, increasing leakage currents
  - Oxygen contaminates BCB and prolonged exposure will compromise the cure and ruin the sample
- Prepare the ‘Blue Oven’ – run N<sub>2</sub> through chamber at 100%
- Prepare ozone reactor – run empty, 20 min
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120 °C, 10 min
- Cool wafer, 5 min
- Surface preparation – clean wafer surface in ozone reactor, 10 min
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, ***NO WATER RINSE***
- Coat wafer with BCB 3022-35 – let sit on surface for 30 sec
- Spin BCB 1.5 kRPM, 30 sec – BCB thickness  $\approx 1.88 \mu\text{m}$
- Place sample into Al cup holder
  - The holder should be deformed such that the sample is completely level, yet has minimal contact with the bottom holder surface
  - Otherwise, cured BCB that has crept onto the bottom surface may prevent the sample from being removed from the holder
- Place sample (in holder) into the ‘Blue Oven’
- Reduce N<sub>2</sub> flow to 60% after 3 min
- Load and run Program 5 (see passivation step)
- Remove sample and inspect under the microscope
- Turn off the ‘Blue Oven’
- Clean carrier wafer in Panasonic ICP – CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 7 min

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- (a) Load sample on carrier wafer, BCB ICP etch  $\text{CF}_4/\text{O}_2$  50:200 sccm, 90 sec
  - (b) Inspect sample in FEI SEM to see if the interconnect posts are exposed
  - Repeat the above etch (45 sec increments) and inspection until all contact and posts are exposed
    - Be sure to run the wafer clean program in between each etch cycle
46. **Metal-2 to Metal-3 interconnect post lithography – 2**
- Repeat this step
47. **Metal-2 to Metal-3 interconnect post deposition – 2**
- Repeat this step
48. **Thin-film BCB microstrip wiring environment – layer 2**
- Repeat this step
49. **Metal-3 adhesion layer**
- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
  - Dehydration bake 120 °C, 10 min
  - Cool wafer, 5 min
  - Prepare ozone reactor – run empty, 20 min
  - Surface preparation – clean wafer surface in ozone reactor, 10 min
  - Surface preparation –  $\text{NH}_4\text{OH}:\text{DI water}$  1:10 dip 10 sec,  $\text{N}_2$  dry, ***NO WATER RINSE***
  - Deposit 100 nm  $\text{SiN}_x$  on the BCB surface by PECVD
  - Photoresist spin – SPR-510, 4 kRPM, 30 sec
  - Pre-exposure PR bake – 90 °C, 60 sec
  - Shoot ‘Metal-3 via’ pattern in stepper, 2.0 sec
  - Post-exposure PR bake – 110 °C, 60 sec
  - Development – MF-701 developer, 90 sec
  - Rinse wafer – DI water for 2 min,  $\text{N}_2$  dry
  - Inspect wafer using optical microscope

## APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW

- Clean carrier wafer in Panasonic ICP – CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 7 min
- Load sample on carrier wafer for ICP etching
  - (a) SiN<sub>x</sub> ICP etch CF<sub>4</sub> 125 sccm, 90 sec
  - (b) Short BCB ashing CF<sub>4</sub>/O<sub>2</sub> 50:200 sccm, 10 sec
- Inspect sample in FEI SEM to see if the interconnect posts are exposed
- Strip photoresist mask
  - (a) Flood expose sample, 15 sec
  - (b) Develop exposed PR – MF-701, 2 min
  - (c) Strip remaining PR in acetone – 3 min
  - (d) Inspect – if PR remaining, use 1165 stripper, 80 °C, 3 min
  - (e) Gently agitate sample surface with small pipet
- Rinse sample in 2-propanol before transferring to DI water or Solvent clean

### 50. Metal-3 interconnect and ground plane lithography

- Solvent clean – 3 min acetone, 3 min 2-propanol, 3 min DI water rinse
- Dehydration bake – 120 °C, 10 min
- Photoresist spin – nLOF 2020, 3 kRPM, 30 sec
- Pre-exposure PR bake – 111°C, 60 sec
- Shoot ‘Metal 3’ pattern in stepper – 0.47 sec
- Post-exposure PR bake – 114°C, 60 sec
- Development – MF-701 developer, 2 min
- Rinse wafer – DI water for 2 min, N<sub>2</sub> dry
- Inspect wafer using optical microscope

### 51. Metal-3 interconnect and ground plane deposition

- Prepare ozone reactor – run empty, 20 min
- Surface preparation – clean wafer surface in ozone reactor, 5 min
- Vent E-beam 4, load private sources – Ti and Au
- Surface preparation – NH<sub>4</sub>OH:DI water 1:10 dip 10 sec, N<sub>2</sub> dry, **NO WATER RINSE**
- Load sample in E-beam 4 – orient long-axis of emitter perpendicular to the direction of sample rotation inside E-beam 4

*APPENDIX B. IMPLANTED COLLECTOR INP HBT / CIRCUIT PROCESS FLOW*

- Allow system to pump-down for 90 min to  $< 10^{-6}$  torr
- Deposit Metal-3 (ground plane)
  - Ti 100 Å (1 Å/sec)
  - Au 15 kÅ
    - \* 2 Å/sec for 1-300 Å
    - \* 3 Å/sec for 301-500 Å
    - \* 4 Å/sec for 501-1000 Å
    - \* 5-6 Å/sec 1001-15000 Å
  - Ti 100 Å (1 Å/sec)
- Metal liftoff – 1165 stripper, 80 °C, 2 hour
- Remove metal and place sample in fresh 1165 stripper, 80 °C, 20 min
  - Gently agitate sample surface with small pipet
  - Rinse sample in 2-propanol before transferring to DI water or Solvent clean